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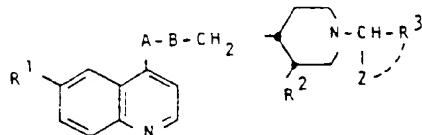
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(54) Quinoline derivatives, processes for their preparation, their use, pharmaceutical compositions containing them and a method for the preparation of these pharmaceutical compositions.

(57) The invention is concerned with novel quinicine and cinchonine derivatives having cardiovascular activities of the formula or a salt thereof,



EP 0 030 044 A1
 in which A-B is -CH₂-CH₂, -CHOH-CH₂-, -CH₂-CHOH-, -C(O)-CH₂-, -CH₂-C(O)-, -C(NOR')-CH₂- or -CH₂-C(NOR')-; R¹ is hydrogen, hydroxy or lower alkoxy; R² is ethyl or vinyl; R³ is C₂-6 alkyl, C₁-6 hydroxylalkyl, lower alkoxy-alkyl or lower alkanoyloxyalkyl, C₃-6 cycloalkyl, hydroxycycloalkyl, lower alkoxycycloalkyl or lower alkanoyloxycycloalkyl, cycloalkyl lower alkyl, hydroxy-, lower alkoxy- or lower alkanoyloxycycloalkyl lower alkyl; cyano, cyano lower alkyl, lower alkenyl, lower alkynyl, tetrahydrofuryl, mono- or di-lower alkylamino

lower alkyl, mono- or di-lower alkylamino lower hydroxy alkyl; optionally substituted phenyl, phenyl lower alkyl or phenyl hydroxy lower alkyl, optionally substituted diphenyl lower alkyl, optionally substituted phenyl lower alkenyl, optionally substituted benzoyl or benzoyl lower alkyl, optionally substituted heteroaryl or heteroaryl lower alkyl, or optionally substituted heteroaryl or heteroaroyl lower alkyl, R¹ is lower alkyl, and Z is hydrogen, lower alkyl or optionally substituted phenyl, or Z and R³ together with the carbon atom to which they are attached form a C₃-6 cycloalkyl group, whereby the substituents at the 3- and 4-position of the piperidine ring are in the cis-position, excluding N-[C₂-6 alkyl, C₂-6 hydroxylalkyl, NN-di-lower alkylamino lower alkyl, optionally substituted C₇-11 alalkyl] substituted derivatives of quinicine and cinchonine.

The compounds of the formula may be in the form of their optically active enantiomers and/or their therapeutically acceptable salts.

Methods for the preparation of the compounds of the formula are also disclosed and form part of the invention.

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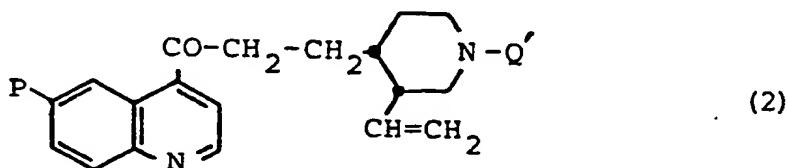
See front page - 1 -

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NOVEL QUINOLINE DERIVATIVES, PHARMACEUTICAL COMPOSITIONS CONTAINING SUCH COMPOUNDS, AND METHOD FOR THE PREPARATION OF THESE COMPOUNDS.

5 The invention relates to novel quinoline derivatives. In French patent publication 2,177,511 and the corresponding German patent application 2,315,148 quinoline derivatives are described with formula 2

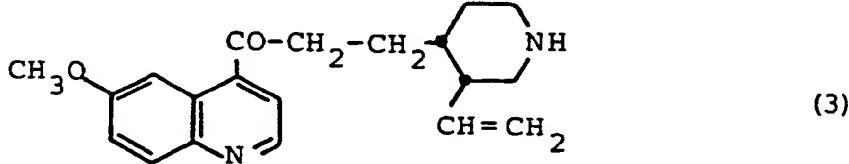
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15 in which P is hydrogen or methoxy and Q' is C_{2-6} alkyl, optionally substituted by alkoxy carbonyl, hydroxy or $-NR'R''$ in which R' and R'' each represent C_{1-4} alkyl or together with the nitrogen to which they are attached form a 5-7 membered heterocyclic ring, which may have oxygen or nitrogen as a
20 second hetero atom, the second nitrogen atom being optionally substituted, R'' may also represent C_{7-11} aralkyl, optionally substituted by halogen, C_{1-6} alkyl or C_{1-6} alkoxy, which compounds have anti-spasmodic and vasodilative activity.

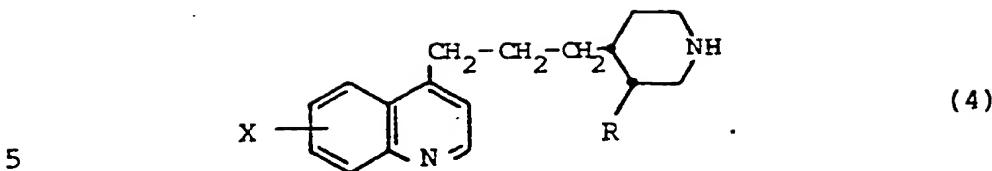
From Ann. Pharm. Fr. 24, 39 (1966) the pharmacodynamic properties of quinicine (formula 3), also named viquidil, are known, in particular in the field of CNS, hypotensive, vasodilative and anti-spasmodic activities.

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In British patent 1,294,538 the use of viquidil in the treatment of cerebral vessel injury, cerebrovascular insufficiency and memory deficiency in humans is described.

In Dutch patent application 77,06614 quinoline derivatives are described with formula 4.

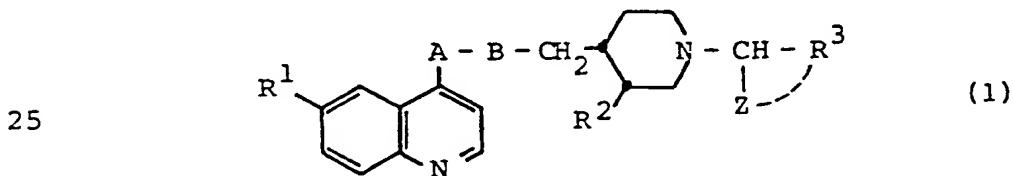


in which R is hydrogen, C₁₋₄ alkyl or C₂₋₄ alkenyl and X is hydrogen, halogen, C₁₋₄ alkyl, alkoxy or alkylthio, trifluoromethyl, nitro, hydroxy, an amino group optionally substituted by one or two C₁₋₄ alkyl groups, or C₁₋₄ acyl or alkylsulphonyl group, which compounds have a serotonin uptake inhibiting effect and anti-arrhythmic activity.

It has now been found, that quinoline derivatives with a substituent at the 4-position containing a N-substituted piperidyl group, possess unexpected pharmacological properties, namely desirable effects on the cardiovascular system, such as anti-hypertensive, anti-thrombotic, vasodilative and anti-arrhythmic activity.

The compounds are particularly useful for use in medicines having anti-hypertensive and anti-arrhythmic activity.

Thus, the invention provides compounds of formula 1,



in which A-B is -CH₂-CH₂-, -CHOH-CH₂-, -CH₂-CHOH-, -C(O)-CH₂-, -CH₂-C(O)-, -C(NOR⁴)-CH₂- or -CH₂-C(NOR⁴)-,
 30 R¹ is hydrogen, hydroxy or lower alkoxy,
 R² is ethyl or vinyl,
 R³ is C₂₋₈ alkyl, C₁₋₈ hydroxyalkyl, lower alkoxyalkyl or lower alkanoyloxyalkyl, C₃₋₆ cycloalkyl, hydroxycycloalkyl, lower alkoxy(cycloalkyl or lower alkanoyloxy)cycloalkyl, cycloalkyl lower alkyl, hydroxy-, lower alkoxy- or lower alkanoyloxy(cycloalkyl) lower alkyl; cyano, cyano
 35

- 3 -

lower alkyl, lower alkenyl, lower alkynyl,
 tetrahydrofuryl, mono- or di-lower alkylamino
 lower alkyl, mono- or di-lower alkylamino lower
 hydroxy alkyl; optionally substituted phenyl,
 5 phenyl lower alkyl or phenyl hydroxy lower alkyl,
 optionally substituted diphenyl lower alkyl,
 optionally substituted phenyl lower alkenyl,
 optionally substituted benzoyl or benzoyl lower alkyl,
 optionally substituted heteroaryl or heteroaryl
 10 lower alkyl, or optionally substituted hetero-
 aryl or heteroaroyl lower alkyl,
 R⁴ is lower alkyl, and
 Z is hydrogen, lower alkyl or optionally substituted
 15 phenyl, or Z and R³ together with the carbon
 atom to which they are attached form a C₃₋₆
 cycloalkyl group,
 whereby the substituents at the 3- and 4-position of the pipe-
 ridine ring are in the cis-position, excluding N-/C₂₋₆ alkyl,
 C₂₋₆ hydroxyalkyl, NN-di-lower alkylamino lower alkyl, option-
 20 ally substituted C₇₋₁₁ aralkyl/ substituted derivatives of
 quinicine and cinchonicine.

As is usual, the carbon chains of the different groups
 may be straight or branched.

The term "lower" is here used to mean a group with up to
 25 six carbon atoms.

The term "optionally substituted" with respect to phenyl
 includes a phenyl group, which may be optionally substituted
 by one, two or three groups selected from lower alkyl, lower
 alkoxy, halogen or hydroxy (no more than two hydroxy groups).

The term "optionally substituted" with respect to hetero-
 aryl is here used to mean a heteroaryl group, which may be
 substituted by one, two or three groups selected from lower
 alkyl, lower alkoxy or halogen.

Aptly, A-B is -CH₂-CH₂- . Another suitable meaning of A-B
 35 is -CHOH-CH₂- . Also suitable is the meaning of A-B being
 -CH₂-CHOH- . Suitably, A-B is -C(O)-CH₂- . Also suitably, A-B is
 -CH₂-C(O)- . The meaning of A-B being -C(NOR⁴)-CH₂ is also apt,

- 4 -

as well as A-B being $-\text{CH}_2-\text{C}(\text{NOR}^4)-$, R^4 being preferably methyl. It has been found, that the compounds in which A-B is $-\text{CHOH}-\text{CH}_2-$ are preferred compounds in relation to their therapeutic properties.

5 Where R^1 is alkoxy, it is preferably methoxy. R^1 is preferably hydrogen or methoxy. Favourably, R^1 is hydrogen. Also favourably, R^1 is methoxy.

10 R^3 as alkyl is preferably ethyl. Other suitable alkyl groups include n-butyl, iso-butyl, n-pentyl, iso-pentyl, n-hexyl, n-heptyl or n-octyl.

R^3 as cycloalkyl is e.g. cyclopropyl or cyclobutyl.

15 Suitable values for R^3 as hydroxyalkyl, hydroxycycloalkyl or hydroxycycloalkyl lower alkyl include substituents with the formula $-(\text{CH}_2)_n\text{C}(\text{OH})\text{R}^4\text{R}^5$, in which n is 0, 1 or 2, R^4 is hydrogen or C_{1-3} alkyl and R^5 is hydrogen or C_{1-3} alkyl or R^4 and R^5 together with the carbon atom to which they are attached form a C_{3-6} carbocyclic ring. The meaning of 1-hydroxy-1-methyl-ethyl is particularly suitable.

20 Favourable values for R^3 as alkoxyalkyl, alkoxyalkyl or alkoxyalkylalkyl include substituents with the formula $-(\text{CH}_2)_n-\text{C}(\text{OR}^6)\text{R}^4\text{R}^5$ in which R^4 and R^5 are as defined hereinbefore and R^6 is C_{1-3} alkyl and n is 0, 1, 2, 3 or 4.

25 Suitable values for R^3 as alkanoyloxyalkyl, alkanoyloxy-cycloalkyl or alkanoyloxyalkylalkyl include substituents with the formula $-(\text{CH}_2)_n\text{C}(\text{OCOR}^7)\text{R}^4\text{R}^5$, in which n , R^4 and R^5 are as defined above and R^7 is hydrogen or C_{1-3} alkyl.

Where R^3 is lower alkynyl, it is preferably ethynyl.

Where R^3 is lower alkenyl, it is preferably ethenyl.

30 Suitable values for R^3 as mono- or di-lower alkylamino lower alkyl and hydroxyalkyl include substituents with the formula $-(\text{CH}_2)_n\text{NR}^8\text{R}^9$, in which n is 1 or 2, R^8 is hydrogen or C_{1-3} alkyl and R^9 is C_{1-3} alkyl.

35 Suitable values for R^3 as optionally substituted phenyl, phenyl lower alkyl or phenylhydroxy lower alkyl include phenyl, benzyl, α -hydroxy-benzyl or benzyl, optionally substituted by 1, 2 or 3 methoxy groups.

Where R^3 is optionally substituted diphenyl lower alkyl, it is preferably 3-(4,4'-difluorodiphenyl)propyl.

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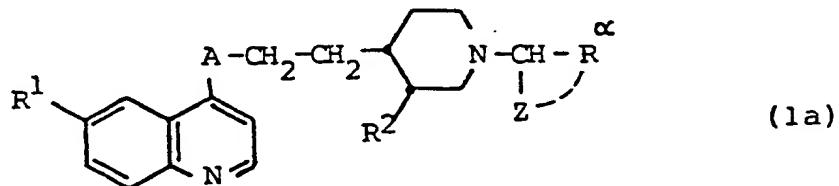
A favourable value for R³ as optionally substituted phenyl lower alkenyl includes cinnamyl.

Suitable values for R³ as optionally substituted benzoyl or benzoyl lower alkyl include benzoyl, fluorobenzoylmethyl, 5 benzoylmethyl, 2-benzoylethyl, 2-(methylbenzoyl)ethyl, 2-(methoxybenzoyl)ethyl, 2-(chlorobenzoyl)ethyl, 2-(fluorobenzoyl)-ethyl and aminobenzoyl. Particularly suitable are the meanings of 2-(4-fluorobenzoyl)ethyl, 2-(4-methylbenzoyl)ethyl and 2-(4-methoxybenzoyl)ethyl and 2-benzoylethyl.

10 Where R^3 is heteroaroyl or heteroaroyl lower alkyl, it is
aptly monocyclic heteroaromatic ring members containing one or
two heteroatoms, of which oxygen, nitrogen or sulfur are pre-
ferred, optionally substituted by a lower alkyl group. Favour-
able values include thiienyl and furyl.

15 Suitable values for Z include hydrogen, methyl, ethyl, n-propyl, isopropyl and phenyl. Preferably, Z is hydrogen.

A particular group of compounds of formula 1 are those of formula 1a



25 in which A is $-\text{CH}_2-$, $-\text{CHOH}-$ or $-\text{CO}-$,
R¹ is hydrogen, hydroxy or lower alkoxy,
R² is ethyl or vinyl,
R^a is C₂₋₈ alkyl, C₁₋₈ hydroxyalkyl, lower alkoxy-
alkyl or lower alkanoyloxyalkyl, C₃₋₆ cycloalkyl, hydroxy-
cycloalkyl, lower alkoxycycloalkyl or lower alkanoyloxy-
cycloalkyl, cycloalkyl lower alkyl, hydroxy-, lower alkoxy-
or lower alkanoyloxcycloalkyl lower alkyl; cyano, cyano
lower alkyl, lower alkenyl, lower alkynyl, tetrahydrofuryl,
mono- or di-lower alkylamino lower alkyl; mono- or di-
loweralkylamino lower hydroxy alkyl; optionally substituted
phenyl, phenyl lower alkyl or phenyl hydroxy lower alkyl,
optionally substituted diphenyl lower alkyl, optionally
substituted phenyl lower alkenyl,

- 6 -

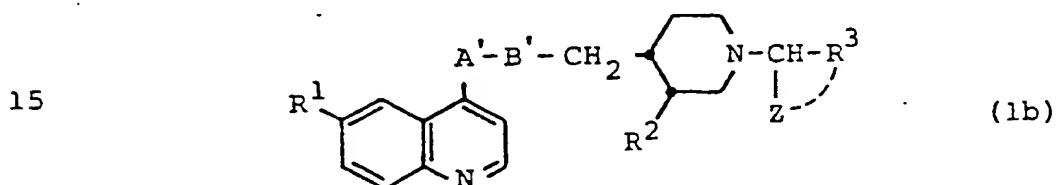
optionally substituted benzoyl or benzoyl lower alkyl, optionally substituted heteroaryl or heteroaryl lower alkyl, or optionally substituted heteroaroyl or heteroaroyl lower alkyl,

5 and

Z is hydrogen, lower alkyl or optionally substituted phenyl, or *Z* and *R*^a together with the carbon atom to which they are attached form a C₃₋₆ cycloalkyl group,

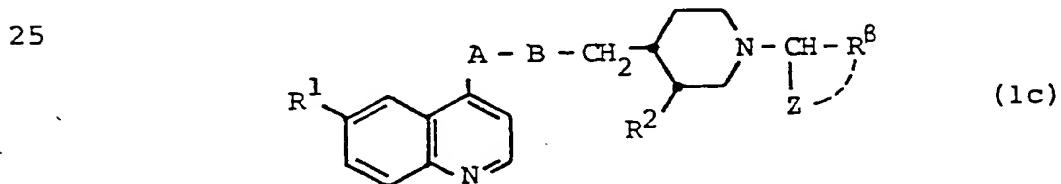
10 excluding the quinicine and cinchonicine derivates as in formula 1.

Another group of particular compounds of formula 1 are those of formula 1b,



in which R^1 , R^2 , R^3 , R^4 and z are as defined above and $A'-B'$
 20 is $-CH_2-CH_2-$, $-CHOH-CH_2-$, $-CH_2-CHOH-$, $-CH_2-C(O)-$, $C(NOR^4)-CH_2-$
 or $-CH_2-C(NOR^4)-$.

A further group of particular compounds of formula 1 are those of formula 1c,

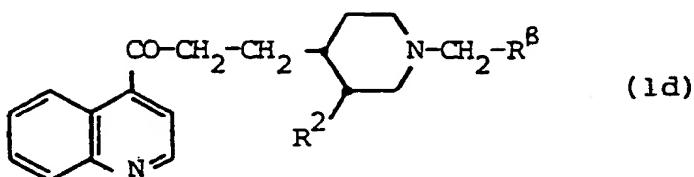


30 in which A-B, R¹, R² and Z are as defined above and
R⁸ is C₁₋₈ lower alkoxyalkyl or lower alkanoyloxy-
alkyl, C₃₋₆ cycloalkyl, hydroxycycloalkyl,
lower alkoxycycloalkyl or lower alkanoyloxycyclo-
alkyl, cycloalkyl lower alkyl, hydroxy-, lower
alkoxy- or lower alkanoyloxycycloalkyl lower
alkyl; cyano, cyano lower alkyl, lower alkenyl,
lower alkynyl, tetrahydrofuryl, mono- or di-

- 7 -

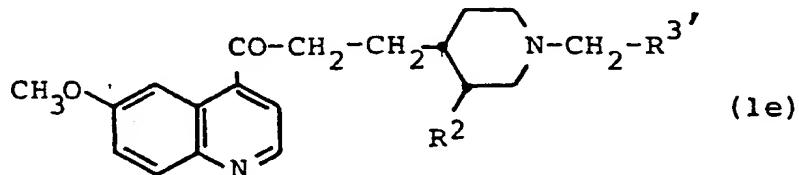
lower alkylamino lower hydroxyalkyl; optionally substituted phenylhydroxy lower alkyl, optionally substituted diphenyl lower alkyl, optionally substituted phenyl lower alkenyl, optionally substituted benzoyl or benzoyl lower alkyl, optionally substituted heteroaryl or heteroaryl lower alkyl, or optionally substituted heteroaroyl or heteroaroyl lower alkyl, or
Z and R^B together with the carbon atom to which they are attached form a C₃₋₆ cycloalkyl group.
Preferably, Z is hydrogen.

A particular group of compounds of formula 1 are those of formula 1d



20 in which R^2 and R^8 are as defined above. In formula 1d we prefer to use R^2 as ethyl. Preferably R^8 is 2-benzoylethyl, 2-(4-fluorobenzoyl)ethyl, 1-hydroxy-1-methyl-ethyl or 3-methoxypropyl.

Another group of preferred compounds of formula 1 are
25 those of formula 1e

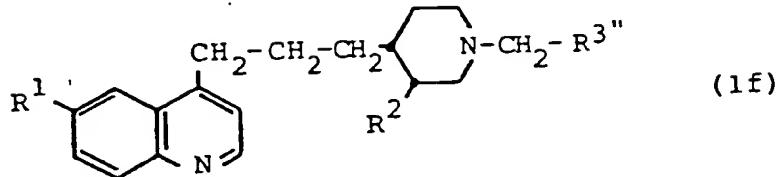


in which R^2 is as defined above, R^3 is ethyl, n-butyl, iso-butyl, n-pentyl, iso-pentyl, lower alkoxy lower alkyl, C_{3-6} cycloalkyl lower alkyl, lower alkenyl, optionally substituted phenyl lower alkyl or optionally substituted benzoyl or benzoyl lower alkyl. R^2 is preferably vinyl. Especially preferred are the compounds of formula Ie in which R^2 is vinyl and R^3 is 2-benzoylethyl, 2-(4-fluorobenzoyl)ethyl or 1-hydroxy-1-methyl-ethyl.

- 8 -

A further group of preferred compounds of formula 1 are those of formula 1f

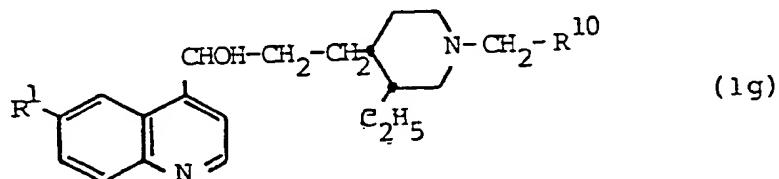
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in which R¹ and R² are as defined hereinbefore and R^{3''} is ethyl, n-butyl, iso-butyl, n-pentyl, iso-pentyl, cyclobutyl, lower alkenyl, optionally substituted phenyl lower alkyl or optionally substituted benzoyl or benzoyl lower alkyl. Especially apt are the compounds in which R¹ is methoxy, R² is vinyl and R^{3''} is cyclobutyl, 2-benzoylethyl or 2-(4-fluorobenzoyl)ethyl. Particularly preferred is the compound in which R¹ is hydrogen, R² is ethyl and R³ is 2-benzoylethyl.

A particular group of compounds of formula 1 are those of formula 1g

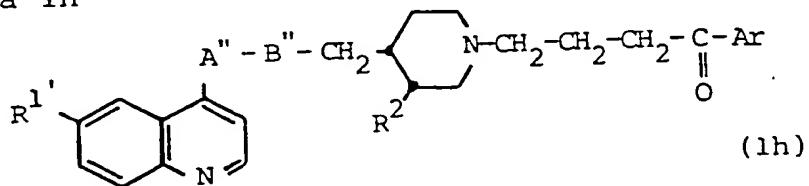
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in which R¹ is as defined above and R¹⁰ is alkyl or -(CH₂)_n-C(O)R¹¹, wherein R¹¹ is an optionally substituted phenyl group and n is 0, 1, 2, 3 or 4. Preferably, R¹⁰ is ethyl, n-propyl or n-butyl, while R¹ is hydrogen or methoxy, or R¹⁰ is 2-benzoylethyl, while R¹ is hydrogen.

Particularly suitable anti-hypertensive agents are those of formula 1h

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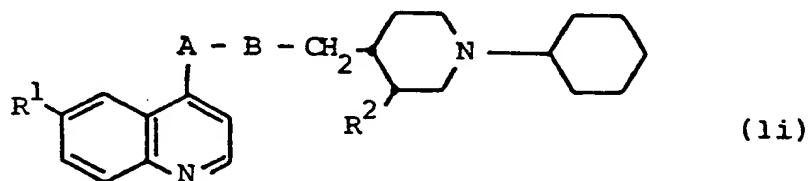


- 9 -

in which A''-B'' is $-\text{CHOH}-\text{CH}_2-$ or $-\text{CH}_2-\text{CHOH}-$, R¹ is hydrogen or methoxy, R² is ethyl or vinyl and Ar is phenyl, thienyl or phenyl substituted by one, two or three groups, selected from fluorine, chlorine or methoxy. Especially suitable values for
 5 Ar include phenyl, fluorophenyl, chlorophenyl, methoxyphenyl, and thienyl. Most suitably A''-B'' is $-\text{CHOH}-\text{CH}_2-$. Most suitably R¹ is hydrogen. Most suitably R² is ethyl. Preferably Ar is phenyl or 4-fluorophenyl, of which 4-fluorophenyl is especially preferred. From the foregoing it will be realized that a
 10 particularly desirable anti-hypertensive compound of this invention is that of formula 1e, in which A''-B'' is $-\text{CHOH}-\text{CH}_2-$, R¹ is hydrogen, R² is ethyl and Ar is 4-fluorophenyl or phenyl.

Other particularly suitable anti-hypertensive agents are those of formula 1i

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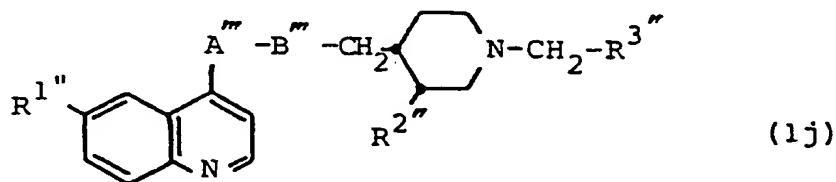


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in which A-B, R¹ and R² are as defined above. Preferably, A-B is $-\text{CH}_2-\text{CH}_2-$ or $-\text{CHOH}-\text{CH}_2-$ or $-\text{C}(\text{O})-\text{CH}_2-$.

Particularly suitable anti-arrhythmic compounds are those of formula 1j

25



30

in which A'''-B''' is $-\text{CHOH}-\text{CH}_2-$, $-\text{CH}_2-\text{CHOH}-$ or $-\text{CH}_2-\text{CH}_2-$, R^{1''} is hydrogen or methoxy, R^{2''} is ethyl or vinyl and R^{3''} is alkyl, hydroxy alkyl or alkoxyalkyl. Most aptly A'''-B''' is $-\text{CHOH}-\text{CH}_2-$. Most aptly R^{1''} is methoxy. Most suitably R^{2''} is ethyl. Most suitably R^{3''} is ethyl. From the foregoing it will be realized that a particularly desirable anti-arrhythmic compound of this invention is that of formula 1j, in which R^{1''}

- 10 -

is methoxy and R^{2''} is ethyl.

From the foregoing it will be appreciated that the following compounds according to the invention are particularly preferred:

- 5 a. N-(3-benzoyl)propyl-hydrocinchonicine
- b. N-(3-benzoyl)propyl-hydrocinchonicinol-1
- c. N-(3-benzoyl)propyl-desoxo-hydrocinchonicine
- d. N-propyl-hydroquinicinol-1
- e. N-butyl-hydroquinicinol-1
- 10 f. N-pentyl-hydroquinicinol-1
- g. N-pentyl-hydrocinchonicinol-1
- h. N-(4-methoxy)butyl-hydrocinchonicine.

The preceding compounds of formula 1 may exist in free base form or in the form of their acid addition or quaternary ammonium salts, for example their salts with mineral acids, e.g. hydrochloric acid, hydrobromic acid or sulphuric acid, or organic acids e.g. acetic acid, fumaric acid or tartaric acid. Naturally the acid used will be pharmaceutically acceptable.

The compounds of formula 1 in which A or B is -CHOH- contain an asymmetric carbon atom and therefore two stereoisomers may exist, provided that there are no asymmetric carbon atoms in a side chain. One or more asymmetric carbon atoms in the N-substituent may give rise to several diastereoisomeric forms.

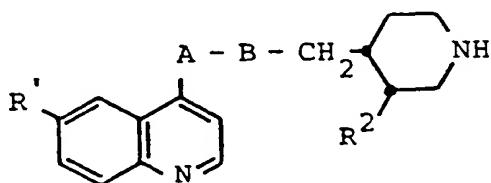
The compounds of the invention are obtainable in crystalline form. They may also be obtained in the form of solvates such as hydrates.

The compounds of the invention, as represented by formula 1, include free base, acid addition and quaternary ammonium salt forms, racemates, separated optical forms and mixtures thereof.

The invention also provides a process for the preparation of compounds of formula 1 in which

A) a compound of formula 5,

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(5)

- 11 -

in which A-B, R¹ and R² are as defined above, is alkylated with a compound of formula 6,

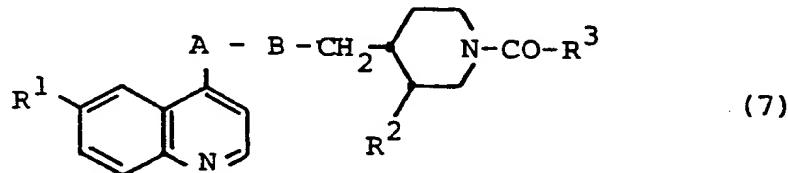
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in which R³ and Z are as defined above and Y' is a nucleophilic
10 leaving group, particularly chlorine, bromine, iodine, aryl-, aralkyl- or alkylsulphonyloxy, and especially mesyloxy or tosyloxy,

or B) a compound of formula 7,

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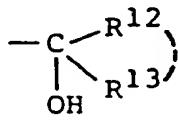


20 in which A-B, R¹ and R² are as defined above, is reduced, to give a compound of formula 1 in which Z is hydrogen,
or C) a compound of formula 5 above is reacted with an epoxide of formula 8,

25



30 in which R¹² is C₁₋₆ alkyl, lower alkenyl, lower alkynyl, optionally substituted phenyl and phenyl lower alkyl and R¹³ is hydrogen or lower alkyl or R¹² and R¹³, together with the carbon atom to which they are attached, form a C₃₋₆ cycloalkyl group, to give a compound of formula 1 in which Z is hydrogen
35 and R³ is



as defined above,

- 12 -

- or D) a compound of formula 1 in which R³ contains a hydroxy group is alkylated,
- or E) a compound of formula 1 in which R³ contains a hydroxy group is acylated,
- 5 or F) a compound of formula 5 above is reacted with a compound of formula 9,



- 10 in which R³ and Z are as defined above, in the presence of a reducing agent,

- or G) a compound of formula 1, in which A-B contains a carbonyl group is reacted with an O-substituted hydroxylamine derivative of formula YO-NH₂, in which Y is R⁴ or a group replaceable by or convertible into R⁴, R⁴ being as previously defined, whereafter the resulting compound in which Y ≠ R⁴ is converted in a compound in which Y = R⁴,

- 20 or H) a compound of formula 1 in which A-B is -CO-CH₂- or -CH₂-CO- is partially or completely reduced to -CHOH-CH₂- or -CH₂-CHOH-, or -CH₂-CH₂- respectively.

In method A, the reaction is preferably carried out by using an equivalent amount or a small excess of the alkylation agent of formula 6. Suitably an acid binding agent is used which does not react with the alkylating agent. For this purpose sterically hindered amines, e.g. dicyclohexylethyl-amine can be used, but generally inorganic bases such as sodium or potassium carbonate and especially sodium or potassium bicarbonate are preferred.

The reaction is preferably carried out in an inert 30 organic solvent, e.g. acetone, methyl ethyl ketone, methyl isobutyl ketone, tetrahydrofuran, dimethylformamide, dimethylsulphoxide, dioxane, methylene chloride, chloroform, benzene, toluene, xylene or a mixture of such solvents. Methyl ethyl ketone, dimethylformamide and toluene or mixtures thereof are preferred. Generally the reaction may be run from 0°C 35 to the boiling point of the solvent.

With less reactive alkylating agents the reaction can be

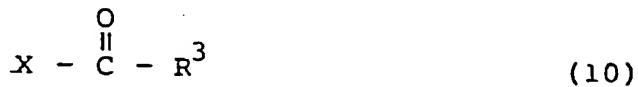
- 13 -

accelerated by addition of catalytic or equivalent amounts of sodium or potassium iodide.

In method B the reduction is suitably carried out using diborane or a complex hydride, such as lithium aluminium hydride. The hydride is added in equivalent amounts or in excess, preferably in quantities up to triple the equivalent amounts. The reduction is preferably carried out in an inert solvent, in particular tetrahydrofuran, at a reaction temperature between 0°C and the boiling point of the solvent.

It must be noted that such reducing agents will also be able to reduce carbonyl or alcohol groups. In compounds of formula 7 in which A or B is -CO- or -CHOH-, these groups usually will be converted into methylene groups leading to compounds of formula 1 in which A or B is reduced to -CH₂-.

The compounds of formula 7 may be prepared for example by acylation of a compound of formula 5 with a compound of formula 10,



in which X is halogen or -OCOR³ and R³ is as defined before.

The acylation of the compounds of formula 5 is preferably carried out in the presence of an acid binding agent, particularly triethylamine or pyridine. Suitable solvents include chloroform, pyridine or dimethyl formamide. Usually the reaction temperature is between 0°C and the boiling point of the reaction mixture.

The compounds of formula 7 may also be prepared by reaction of a compound of formula 5 with a carboxylic acid of formula 10 (X is OH), in which R³ is as defined above, in the presence of dicyclohexylcarbodiimide. This acylation method is effected under normal conditions, e.g. in chloroform as a solvent, after which the reaction product is isolated in a conventional manner.

Method B is advantageous for the preparation of compounds of formula 1 in which R³ contains a branched alkoxyalkyl group.

Method C is suitably carried out in an inert organic

- 14 -

solvent, preferably a lower alcohol of 1-5 carbon atoms or in a mixture of such an alcohol with dichloromethane. Sometimes it is advantageous to add water to the reaction mixture.

The reaction conditions usually depend on the reactivity 5 of the epoxide. Usually the reaction may be run some hours and is preferably effected at temperatures in the range of 20-120°C. Where a volatile epoxide is used, a closed system may be necessary.

In method D, the hydroxy group is suitably first converted 10 into the corresponding alkali salt, e.g. with sodium hydride in an aprotic solvent. This salt is then treated with an alkyl halide or an alkyl or aryl sulphonic ester, preferably with an alkyl halide.

Method E is preferably carried out with an acide chloride 15 or anhydride, as described under method B.

Method F is suitably carried out with hydrogen as reducing agent in the presence of a catalyst, for example palladium on coal. Depending on the catalyst of choice hydrogen pressures of 1-150 atm are involved. The reaction conditions 20 are as commonly used for this type of reaction. The reaction is suitably carried out in a solvent, such as a lower alcohol, preferably methanol or ethanol at a temperature generally in the range of 20-100°C, preferably between 20-40°C.

Another suitable method in which the use of high pressure 25 is avoided comprises the reaction of the compounds of formulae 5 and 9 in the presence of an equivalent amount of sodium cyanoborohydride and a base, such as potassium hydroxide. The reaction is suitably carried out at room temperature, in a solvent such as an alcohol, preferably methanol or ethanol.

Method G is carried out in conventional manner for this 30 type of reaction. Preferably, the reaction is carried out in a solvent, such as an alcohol, dioxane, dimethyl formamide, tetrahydrofuran or pyridine, at a temperature generally between room temperature and the boiling point of the reaction mixture. 35 The hydroxylamine derivative is usually added as an acid salt, preferably the hydrochloride, which salt is preferably dissolved in pyridine. Sometimes it may be advantageous to convert

- 15 -

the starting carbonyl compound first to, for example, the corresponding oxime. This oxime may then be converted to its sodium salt, which may be easily alkylated to the desired product.

5 It will be appreciated by those skilled in the art, that the conversion of the carbonyl group to the oxime ether group may occur with both the carbonyl compounds of formula 1 and formula 5.

Method H, the partial or complete reduction of a compound of formula 1, in which A-B is $-\text{CO}-\text{CH}_2-$ or $-\text{CH}_2-\text{CO}-$ may 10 be carried out in conventional manner. A suitable reducing agent for the conversion to the desoxo compound ($-\text{CH}_2-\text{CH}_2-$) is e.g. hydrazine hydrate, in the presence of an alkali metal hydroxide, such as potassium hydroxide, in a suitable solvent 15 such as an alcohol, e.g. ethylene glycol. A suitable reducing agent for the partial reduction to the alcohol derivative ($-\text{CHOH}-\text{CH}_2-$) is for example a complex hydride, such as sodium borohydride. This reduction is advantageously carried out at a temperature of about -5 to -10°C in a suitable solvent, like 20 an alcohol, preferably methanol, ethanol or isopropylalcohol. If desired, the alcohol compound may also be converted into the corresponding desoxo compound, e.g. by converting the alcohol in a suitable solvent, such as ethylalcohol, with phosphorous pentachloride to the chloride and reducing the 25 resulting compound, for example with hydrogen gas in a solvent, such as ethylalcohol and for example palladium on coal as a catalyst.

The starting materials of formula 5 are either known or may be prepared in conventional manner from known compounds.

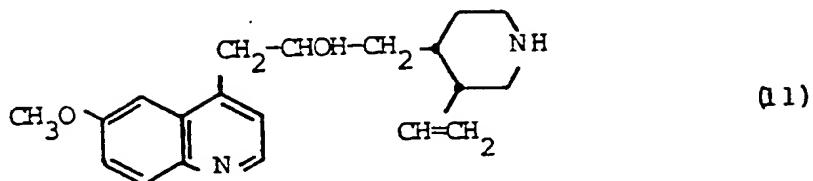
30 Compounds of formula 5, in which A-B is $-\text{C}(\text{O})-\text{CH}_2-$ may be prepared according to the methods described in French patent 73,41043 (publ. nr. 2,206,944) and in Dutch patent application no. 77,06614.

The methods described in said patents are based on 35 the condensation of an ester of 3-(4-piperidyl)propionic acid with a quinoline derivative, which is substituted at the 4-position by a carboxylic ester group or a lithium atom.

- 16 -

In J. Amer. Chem. Soc. 100, 576-581 (1978) the preparation of a compound of formula 11

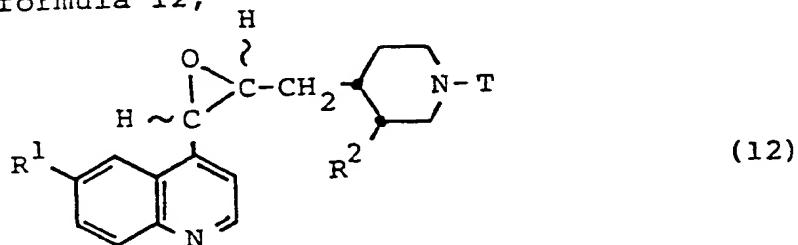
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is described by converting 6-methoxylepidine in situ to 6-methoxylepidyllithium and reacting this compound with the methyl ester of a (4-piperidyl) acetic acid derivative (N-benzoylmeroquinene methyl ester). While removing the N-benzoyl group of the resulting keto compound of the 1,3-disubstituted propanone-2 type with DIBA, the compound is reduced to a 15 propanol-2 derivative.

Compounds of formula 5 in which A-B is $-\text{CH}_2\text{-CHOH}-$, may also be prepared e.g. by reduction of a cis- or trans-oxirane compound of formula 12,

20

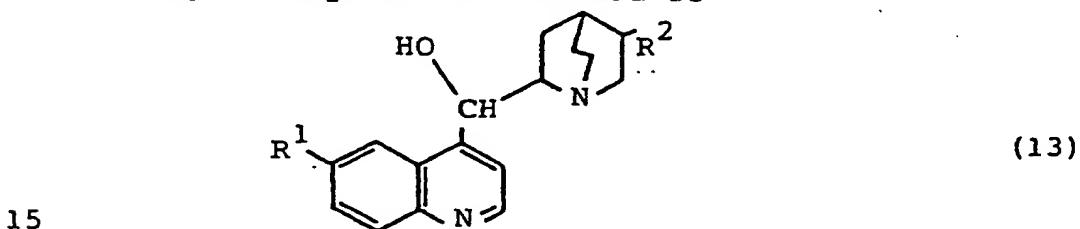


25 in which R^1 and R^2 are as defined above and T is a protecting group and preferably benzyl, followed by removal of this protecting group in conventional manner. The reduction is suitably carried out by leading hydrogen gas through a suitable solvent, such as an alcohol, e.g. ethylalcohol, in the presence of a suitable catalyst, e.g. palladium on coal, at room temperature or slightly elevated temperature. As a result of the reduction generally alcohols are formed as a mixture of diastereoisomers, which may be separated in conventional manner. The removal of the protecting group may be carried out with known techniques. If the protecting group is alkyl, this group may be removed e.g. with cyanogen bromide or chlorocarbonic acid ester. If the protecting group is a benzyl group, debenzylation occurs preferably catalytically.

- 17 -

It is noted, that the protecting group T may have the meaning of $-\text{CH}_2\text{R}^3-$, which is previously defined. In that case the reduction will result into a compound with formula 1, so that the removal of the protecting group may be omitted.

5 The preparation of the cis- and trans-oxirane compounds of formula 12 have been described by L. Keefer, Thesis Univ. of Hampshire 1966 and G.G. Lyle and L.K. Keefer, Tetrahedron 23, 3253-3263 (1967) or may be prepared in an analogous way. Generally, the compounds may suitably be prepared by quaternizing a compound of formula 13



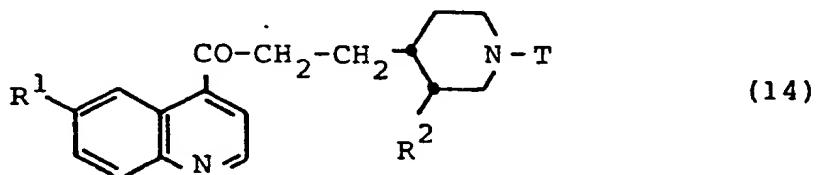
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in conventional manner, for example to the corresponding benzobromide and converting the resulting compound with a base.

Because of the stereospecificity of the reaction a compound 20 of formula 13 in the erythro configuration is preferably used as the starting material, while the quaternizing group is not too small, i.e. larger than methyl and ethyl. Thus, a suitable group is for example benzyl. The reaction with the quaternizing compound is suitably carried out with a base, such as potassium 25 hydroxide in a solvent, such as ethylalcohol.

It is noted, that if the above-described reaction is carried out with the quaternary salt of a threo compound of formula 13, a keto compound of formula 14 may be formed,

30



35 in which R¹, R² and T are as previously defined. If a relatively small quaternizing group is used, such as a methyl group, generally a keto compound of formula 14 is formed in this

reaction, both if a threo or an erythro compound of formula 9 is the starting material. Therefore, this method is also suitable for the preparation of compounds of formula 1, in which A-B is $-\text{CO}-\text{CH}_2-$ and, after removal of the protecting group T,
5 of the corresponding compounds of formula 5.

Threo compounds of formula 13 may also be converted to oxirane compounds of formula 12, if the quaternizing group is not too small, e.g. benzyl. This reaction is carried out with a strong base, in which B⁻ is bulky, for example potassium t-
10 butoxide in t-butanol. The resulting oxirane compound is usually in the cis-configuration.

The resulting compounds of formula 1 or formula 5, in which R¹ and R² are as previously defined and A-B is $-\text{CH}_2-\text{CHOH}-$,
15 may be oxidized in conventional manner to the corresponding keto compounds, in which A-B is $-\text{CH}_2-\text{CO}-$. A suitable method includes the Oppenauer oxidation. Such keto compounds may also be prepared by the cited method described in J. Amer. Chem. Soc. 100, 576-581 (1978), for example by condensing 4-methyl-
20 quinoline which is optionally substituted at the 6-position, with the ester of a 4-piperidyl-acetic acid derivative under the influence of lithium and a strong base.

The reaction products from any method A-H may be isolated from the reaction mixture and purified by conventional means.

In a number of cases, certain reaction steps may be
25 carried out in a different sequence or simultaneously or without isolating intermediates, and these possibilities are all included in the invention

Those skilled in the art will appreciate that protecting groups may be used to protect certain reactive functions during
30 the above processes, in accordance with conventional chemical practice.

Certain compounds of formula 1 may also be used for the preparation of other compounds of formula 1 and are therefore also suitable as intermediates.

35 Diastereoisomers may be separated by known techniques,

based on their different physical and chemical characteristics, e.g. by fractional crystallisation or by column chromatography. These isomer separations may be effected after the final step of the synthesis used or optionally at a previous stage, after 5 the formation of the mixture of diastereoisomers.

Racemic mixtures may be resolved into their enantiomers, in conventional manner, e.g. by separation of their salts with suitable optically active acids.

The free base and acid addition salt forms of the com-10 pounds of formula 1 may be interconverted by standard methods.

The quaternary compounds can be made by refluxing the N-substituted compounds having the desired substituent onto the piperidine-nitrogen in the presence of suitable alkyl halides. Suitable solvents for quaternisation include methanol, ethanol, 15 tetrahydrofuran or glycerol alone or in combination with minor amounts of water, reaction temperatures varying from room temperature to the boiling point of the solvent or mixture used. Other wellknown methods such as the use of lithiumalkyl com-20 pounds or alkylmagnesium halides may be also applied. All different quaternary compounds can be made except those in which steric hindrance of the already present piperidine-25 substituent inhibits the formation of quaternary salts.

The compounds of formula 1 possess pharmacological activity. In particular they possess cardiovascular activity, 25 for example anti-hypertensive, anti-thrombotic, vasodilative and anti-arrhythmic activity.

An indicated suitable daily dosage (for a 70 kg human) is from 1 to 200 mg, of a compound of formula 1, preferably administered in divided dosages of from 0.5 mg to 50 mg 2 to 30 4 times daily, or in retard form.Orally administrable unit dose forms may thus contain 0.5, 1, 2.5, 5, 10, 20, 25 or 50 mg of an active ingredient.

The compounds may be administered in free base form or in the form of their pharmaceutically acceptable acid addition 35 salt forms, which salt forms have the same order of activity as the free base forms.

- 20 -

The compounds of formula 1 may be admixed with conventional pharmaceutically acceptable diluents or carriers and, optionally, other excipients, and administered for example in such forms as tablets, capsules and injectable 5 solutions. They may also be administered in combination preparations with other active agents.

The pharmaceutical compositions may be formulated in conventional manner, e.g. as for other anti-hypertensive agents.

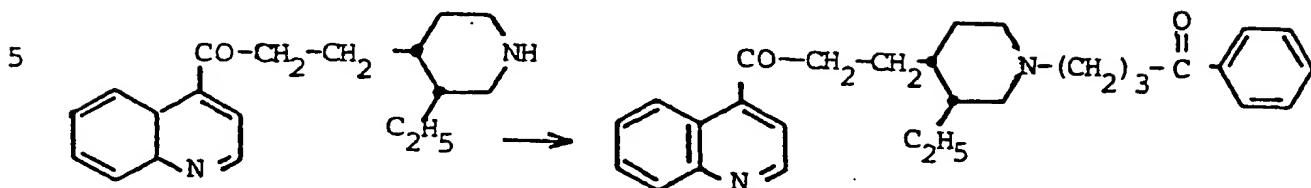
10 The invention also relates to a method for the preparation of a pharmaceutical composition, characterized in that at least a compound of formula 1, as defined hereinbefore, or a pharmaceutically acceptable salt thereof, is brought in a form suitable for therapeutic purposes.

15 The invention further relates to a method of treating mammals, in particular humans, suffering from e.g. cardiovascular diseases, which comprises administering an effective amount of a compound of formula 1 as defined hereinbefore or a pharmaceutically acceptable salt thereof, preferably in the 20 form of a pharmaceutical composition.

The following Examples illustrate the invention.

Example 1

N-(3-benzoyl)-n-propyl-hydrocinchonicine



A solution of 249.3 g (0.75 mole) of hydrocinchonicine
10 HCl in 2000 ml of water was made alkaline with 4N sodium
hydroxide (pH 9-10) and extracted with 1000 ml of toluene.
The water phase was separated from the toluene and the
toluene phase was dried over mol. sieves (4 Å) and filtered.

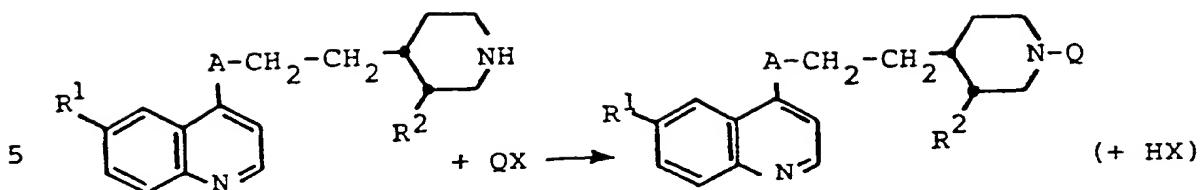
To the dried solution 207 g (1.5 mole) of potassium
15 carbonate and 164.3 g (0.9 mole) of γ -chlorobutyrophenone
were added, and the mixture was refluxed for 24 hours with
stirring. A precipitate of potassium chloride was formed
during the reaction. The conversion was followed by thin-
layer chromatography (silica gel with chloroform/acetone/
20 diethyl amine 5:4:1 as the eluent). After the reaction
mixture was cooled to room temperature water was added, the
layers were separated and the toluene phase concentrated in
vacuo to about 250 ml. The solution was filtered over a
short silica gel column, chloroform being the eluent. The
25 fractions with the desired product were evaporated to dry-
ness in vacuo and the title compound remained as an oil.

Example 2

N-(3-benzoyl)-n-propyl-hydrocinchonicine bifumarate

The oil obtained according to Example 1 was dissolved
in acetone. Fumaric acid was then added to the boiling solu-
tion till pH 6-7 was reached. Upon cooling to room tempera-
ture, the title compound crystallized. The salt was recrys-
tallized from acetone/methanol, melting point 127-129°C.

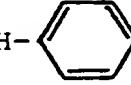
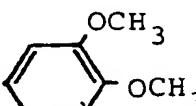
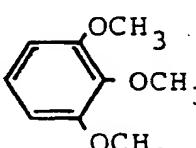
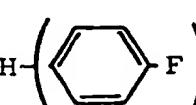
In the same way the following compounds were prepared:

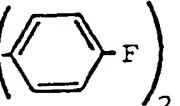
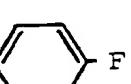
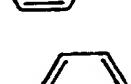


Ex	A	R ¹	R ²	Q	m.p. °C/salt	X		
10	1 (rep.)	CO	H	ethyl	(CH ₂) ₃ -CO-		(oil)	Cl
10	2 (rep.)	CO	H	ethyl	(CH ₂) ₃ -CO-		127-129 BF	Cl
15	3	CO	OCH ₃	vinyl	(CH ₂) ₃ -CO-		117-119 BF	Cl
15	4	CHOH	H	ethyl	(CH ₂) ₃ -CO-		184-186 BO	Cl
20	5	CHOH	OCH ₃	ethyl	(CH ₂) ₃ -CO-		148-151 BO	Cl
25	6	CH ₂	H	vinyl	(CH ₂) ₃ -CO-		119-121 BF	Cl
25	7	CH ₂	H	ethyl	(CH ₂) ₃ -CO-		191-192 BO	Cl
30	8	CH ₂	OCH ₃	vinyl	(CH ₂) ₃ -CO-		92-94 HCl	Cl
30	9	CO	H	ethyl	(CH ₂) ₃ -CO-		121-123 BF	Cl
35	10	CH ₂	H	vinyl	(CH ₂) ₃ -CO-		124-126 BF	Cl
35	11	CO	H	ethyl	(CH ₂) ₃ -CO-		175-177 Fu	Cl

Ex	A	R ¹	R ²	Q	m.p. °C/salt	X	
12	CO	H	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{OCH}_3$	139-141 BF	Cl	
5	CH ₂	H	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{OCH}_3$	122-124 BF	Cl	
10	CO	H	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{Cl}$	128-130 BF	Cl	
15	CH ₂	OCH ₃	vinyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{Cl}$	128-131 biHCl	Cl	
15	CO	H	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{F}$	143-146 BF	Cl	
17	CHOH	H	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{F}$	186-188 BO	Cl	
20	CHOH	H	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{F}$	183-185 BO (one isomer)	I	
19	CH ₂	H	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{F}$	196-198 BO	I	
25	CH ₂	OCH ₃	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{F}$	182-183 BO	I	
21	CH ₂	OCH ₃	vinyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{F}$	151-152 BO	I	
30	CHOH	OCH ₃	vinyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_5-\text{F}$	159-161(d) TO(2:3)	I	
23	CO	H	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_6\text{H}_4-\text{Cl}$	122-124 BF	Cl	
35	24	CO	H	ethyl	$-(\text{CH}_2)_3-\text{CO}-\text{C}_4\text{H}_3-\text{Cl}$	157-160 BF	Cl

Ex	A	R ¹	R ²	Q	m.p. °C/salt	X
25	CO	H	ethyl	- (CH ₂) ₂ -CH ₃	187-189 BO	Br
26	CHOH	OH	ethyl	- (CH ₂) ₂ -CH ₃	134-137 TO (2:3)	Br
5	27	CO	OH	ethyl	- (CH ₂) ₂ -CH ₃	158-159 HCl
	28	CO	OCH ₃	ethyl	- (CH ₂) ₂ -CH ₃	202-204 BO
	29	CO	OC ₃ H ₇	ethyl	- (CH ₂) ₂ -CH ₃	135-136 BF
	30	CO	O- iC ₅ H ₁₁	ethyl	- (CH ₂) ₂ -CH ₃	190-192 BO
	31	CHOH	OCH ₃	ethyl	- (CH ₂) ₂ -CH ₃	124 (d) HCl
10	32	CHOH	OCH ₃	ethyl	- (CH ₂) ₂ -CH ₃	136-138 base (one isomer)
	33	CH ₂	H	vinyl	- (CH ₂) ₂ -CH ₃	236-239 biHCl
	34	CH ₂	H	ethyl	- (CH ₂) ₂ -CH ₃	178-180 HCl
	35	CH ₂	OH	ethyl	- (CH ₂) ₂ -CH ₃	176-179 HCl
	36	CH ₂	OCH ₃	ethyl	- (CH ₂) ₂ -CH ₃	145-153 BO
15	37	CH ₂	OC ₃ H ₇	ethyl	- (CH ₂) ₂ -CH ₃	136-138 BO
	38	CH ₂	O- iC ₅ H ₁₁	ethyl	- (CH ₂) ₂ -CH ₃	158-159 BO
	39	CO	H	ethyl	- (CH ₂) ₃ -CH ₃	160-162 BF
	40	CHOH	OCH ₃	ethyl	- (CH ₂) ₃ -CH ₂	115-118 TO (2:3)
	41	CO	H	ethyl	- (CH ₂) ₄ -CH ₃	102-104 BF
20	42	CHOH	H	ethyl	- (CH ₂) ₄ -CH ₃	116-119 BO
	43	CH ₂	H	vinyl	- (CH ₂) ₄ -CH ₃	142-143 BO
	44	CH ₂	H	ethyl	- (CH ₂) ₄ -CH ₃	141-142 BF
	45	CHOH	OCH ₃	ethyl	- (CH ₂) ₄ -CH ₃	106-108 base
	46	CH ₂	OCH ₃	vinyl	- (CH ₂) ₄ -CH ₃	80-83 BO
25	47	CH ₂	OCH ₃	ethyl	- (CH ₂) ₄ -CH ₃	145-147 BO
	48	CO	H	ethyl	- (CH ₂) ₅ -CH ₃	106-108 BF
	49	CO	H	ethyl	- (CH ₂) ₆ -CH ₃	90-93 BF
	50	CO	H	ethyl	- (CH ₂) ₈ -CH ₃	105-107 BF
	51	CO	OCH ₃	vinyl	- (CH ₂) ₈ -CH ₃	134-136 BF
30	52	CH ₂	H	vinyl	- (CH ₂) ₈ -CH ₃	173-174 BO
	53	CH ₂	OH	ethyl	- (CH ₂) ₈ -CH ₃	156-158 BO
	54	CH ₂	OCH ₃	vinyl	- (CH ₂) ₈ -CH ₃	135-137 BO
	55	CO	H	ethyl	- CH ₂ -CH(CH ₃) ₂	148-150 BF
	56	CO	H	ethyl	- (CH ₂) ₂ -CH(CH ₃) ₂	180-182 BF
35	57	CH ₂	OCH ₃	ethyl	- (CH ₂) ₂ -CH(CH ₃) ₂	134 BO
	58	CO	H	ethyl	- (CH ₂) ₃ -CH(CH ₃) ₂	138-140 BF

Ex	A	R ¹	R ²	Q	m.p. °C/salt	X	
59	CO	OCH ₃	vinyl	-CH ₂ -CH ₂ - 	183-185 BF	I	
5	60	CO	H	ethyl	- (CH ₂) ₃ -CH ₂ OH	140-143 BO	Br
61	CO	H	ethyl	-CH ₂ -CH ₂ -(OH)C-(CH ₃) ₂	142-145 BO	Br	
62	CO	H	ethyl	-CH ₂ -CH ₂ -OCH ₃	140-142 BF	Br	
63	CO	OCH ₃	vinyl	-CH ₂ -CH ₂ -OCH ₃	126-128 BF	Br	
10	64	CO	H	ethyl	- (CH ₂) ₃ -O-CH ₃	(oil)	
	65	CHOH	H	ethyl	- (CH ₂) ₄ -OCH ₃	144-148 BO	Br
	66	CO	H	ethyl	- (CH ₂) ₄ -OCH ₃	108-110 BF	Br
	67	CO	H	ethyl	- (CH ₂) ₄ -CN	149-151 BO	Br
15	68	CO	H	ethyl	- CH ₂ - 	201 BO	Br
	69	CO	H	ethyl	-CH ₂ - 	146 (d) BF	Br
20	70	CO	H	ethyl	- (CH ₂) ₂ -N(CH ₃) ₂	(oil)	Cl
	71	CO	H	ethyl	-CH ₂ CHOHCH ₂ -N(C ₂ H ₅) ₂	122-125 TO	Cl
	72	CO	OCH ₃	vinyl	-CH ₂ -CH=CH ₂	97-99 BF	Cl
	73	CH ₂	OCH ₃	vinyl	-CH ₂ -CH=CH ₂	191-193 Pa	Cl
	74	CH ₂	OCH ₃	ethyl	-CH ₂ -CH=CH ₂	145-148 BO	Cl
25	75	CO	H	ethyl	-CH ₂ -CH=CH- 	153-155 BF	Br
	76	CO	H	ethyl	-CH ₂ -C≡CH	203-205 BO	Cl
30	77	CO	H	ethyl	-CH ₂ -CH ₂ - 	170-172 BF	Br
	78	CO	H	ethyl	-CH ₂ -CH ₂ - 	155-160 (d) TO (2:3)	Br
35	79	CO	H	ethyl	- (CH ₂) ₃ -CH- 	170 (d) HCl	Cl

Ex	A	R ¹	R ²	Q	m.p. °C/salt	X
80	CH ₂	OCH ₃	vinyl	- (CH ₂) ₃ -CH- 	124 (d) HCl	Cl
5	CO	OCH ₃	vinyl	-CH ₂ -CO- 	150-152 BO	Br
10	CO	H	ethyl	- (CH ₂) ₂ -CO- 	153-155 BF	Br
15	CO	H	ethyl	- (CH ₂) ₄ -CO- 	175-178 BO	I
	CO	H	ethyl	- (CH ₂) ₃ -CHOH- 	189-192 BO	I

The salts in the Table have been abbreviated as follows:

BF = bifumarate TO = trioxalate

BO = bioxalate

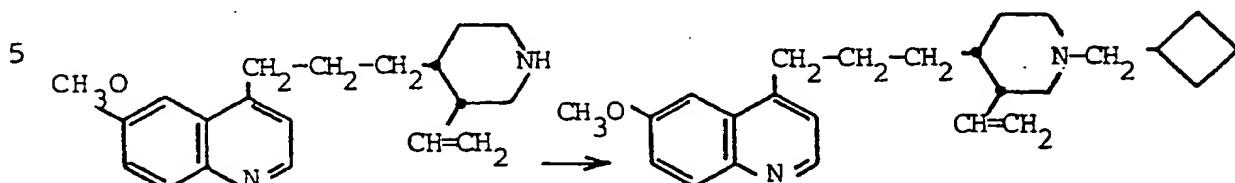
Fu = fumarate

Pa = pamoate

The compounds mentioned in the Table are all Examples
25 which illustrate the invention. For clarity's sake the com-
pounds prepared according to Examples 1 and 2 have also been
mentioned in the Table.

Example 85

N-cyclobutylmethyl-desoxoquinicine hydrochloride



10 To a suspension of 10.0 g (28.8 mmole) of desoxoquinicine.HCl in 125 ml of dichloromethane 58 ml of triethylamine was added with stirring.

In 60 ml of dichloromethane 8.6 g (86.4 mmole) of cyclobutanecarboxylic acid was dissolved, then 10.3 g (86.4 mmole) of thionyl chloride was dropwise added with stirring and the solution was refluxed for 1 h. After cooling to room temperature the resulting solution was dropwise added to the above-mentioned solution of desoxoquinicine in dichloromethane with stirring in a nitrogen atmosphere. The 20 solution was refluxed with stirring for another 2 hours. The reaction mixture was cooled to room temperature and 1000 ml of water was added with stirring.

The layers were separated and the water layer was extracted twice with 50 ml of chloroform. The combined extract of dichloromethane and chloroform was shaken twice with 25 100 ml of water, dried over magnesium sulphate, filtered and evaporated in vacuo. The residue was chromatographed over silica gel with chloroform/acetone/diethylamine 5:4:1 as the eluent. The fractions containing the acid amide were collected and 30 evaporated to dryness in vacuo. The residue was dissolved in 450 ml of dry tetrahydrofuran and dropwise added to a suspension of 3.2 g (86.4 mmole) of lithium aluminium hydride in 120 ml of dry tetrahydrofuran with stirring in a nitrogen atmosphere. Stirring was continued for another 2 hours followed by refluxing for 1 hour.

The solution was cooled to room temperature and 40 ml of ethyl acetate was dropwise added with stirring and cooling in ice water, followed by 300 ml of a 30% solution of ammonium chloride. The organic phase was separated, dried over magne-

sium sulphate, filtered and evaporated to dryness in vacuo.

The residue was chromatographed over silica gel with cyclohexane/acetone 8:1 as the eluent. The fractions containing the N-cyclobutylmethyldesoxoquinicine were evaporated to dryness in vacuo, then the base was converted to the HCl-salt with 1 eq. of isopropylalcohol.HCl. Melting point 140-143°C.

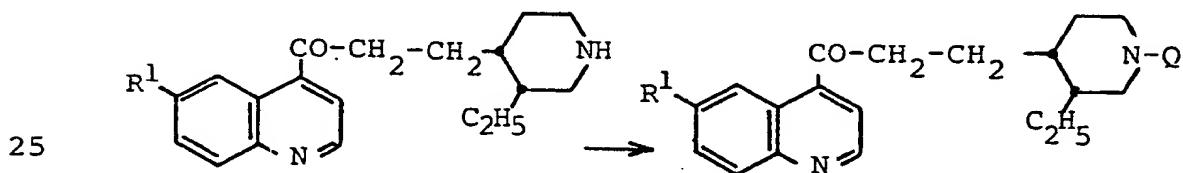
Example 86

N-(2-hydroxy-2-methyl)propyl-hydrocinchonicine bioxalate
(formula: see table below)

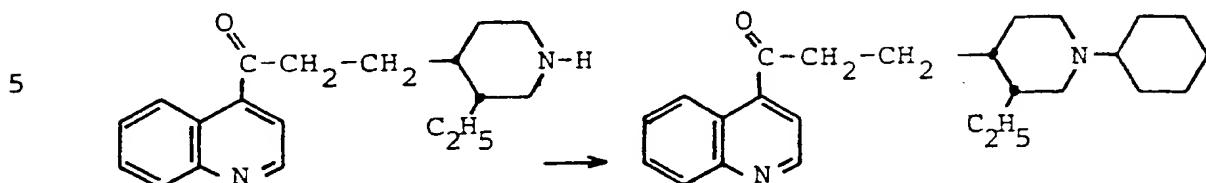
To a solution of 22 g of hydrocinchonicine in 80 ml of
5 absolute alcohol 9 ml of 1,2-epoxy-2-methylpropane was added.
The mixture was refluxed for 2 hours and the conversion was
followed by thin layer chromatography. The reaction mixture
was cooled to room temperature and evaporated to dryness in
vacuo.

10 The residue was purified by preparative high pressure
liquid chromatography with ethyl acetate/diethyl amine 95:5
as the eluent. The fractions containing the purified product
were collected and evaporated to dryness in vacuo. The resi-
due was dissolved in ethyl acetate / methanol. The solution
15 was heated, then the base was converted into the bioxalate
(1:1) with an equivalent amount of oxalic acid. Melting point
132-134°C.

In a similar way the following compounds have been
obtained which are mentioned (next to the compound described
20 in this Example) in the table below:



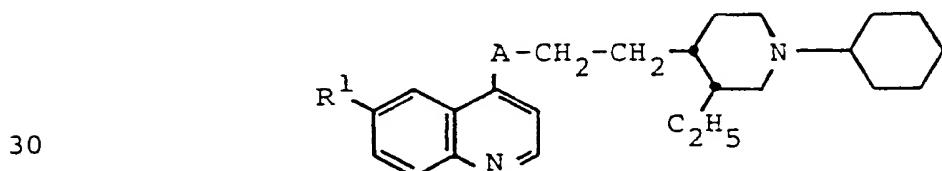
Ex	R¹	Q	m.p. (°C) salt
30	86 (rep.)	-CH₂C(OH)(CH₃)₂	132-134 bioxalate
	87	-CH₂C(OH)(C₂H₅)₂	72-75 bioxalate
	88	-CH₂CH(OH)-	163-165 bioxalate
35	89	-CH₂-	172-174 bioxalate
	90	-CH₂-	82-85 bioxalate
91	H	-CH₂-C(OH)(CH₃)-	228-229 HBr

Example 92N-Cyclohexyl-hydrocinchonicineoxalate

To a solution of 50 ml of methanol with 0.5 g of potassium hydroxide was added with stirring 6.7 g (0.020 mol) of hydrocinchonicinehydrochloride. After stirring for some time 1.96 g (0.020 mol) of cyclohexanone was added. Then 0.44 g (0.007 mol) of sodium cyanoborohydride was added and the reaction mixture was stirred for 30 min. at room temperature. After adding 0.5 g potassium hydroxide stirring was continued for 4h. at room temperature. The reaction mixture was evaporated to dryness in vacuo and treated with ether and water. The ether layer was extracted with 4 n HCl. The water layer was basified with 4 n sodium hydroxide and extracted with ether. The ether extract was dried on magnesium sulfate, filtered and evaporated to dryness in vacuo. The crude product was purified over Silica Gel G with ethylacetate / diethylamine (9:1) as eluent.

With oxalic acid the base was converted to its oxalate (mol. ratio 1:1.25) having a melting point of 108°-109° C.

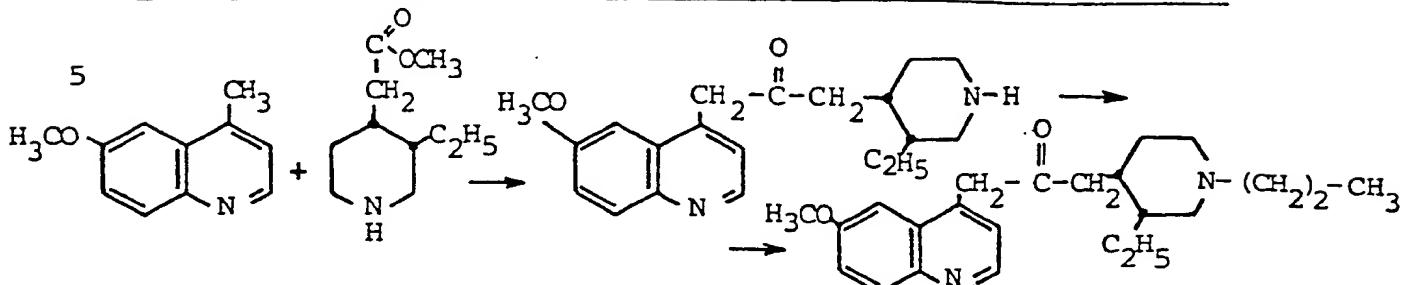
In the same way were prepared:



Ex	A	R¹	m.p. (°C)	salt
92 (rep)	CO	H	108-109	BO
93	CH₂	H	101-104	BO
94	CHOH	H	198-200	BO
95	CH₂	OCH₃	69-70	base

Example 96

1-(6-methoxy-4-quinolyl)-3-[3(R)-ethyl-N-propyl-4-(S)-piperidyl]- propanone-2



10 To 46.6 ml (77.2 mmol) of a 15%-solution of n-butyl lithium in hexane was added under nitrogen 9.2 g (91.1 mmol) of diisopropylamine with stirring at -5° to -10°C over 20 min. Then 11.75 g (67.9 mmol) of 6-methoxylepine in 40 ml of tetrahydrofuran was added dropwise with 15 stirring at -5°C followed by dropwise addition of a solution of 7.6 g (41.1 mmol) of 3(R)-ethyl-4(S)-piperidyl acetic acid methyl ester in 40 ml of tetrahydrofuran. Stirring was continued at -5°C for 3 h.

20 The reaction mixture was acidified (pH 6) with acetic acid and 5 g of potassium bicarbonate was added. Then the reaction mixture was diluted with 100 ml of methanol, filtered and evaporated to dryness in vacuo. To the residue 150 ml of water and 4 n HCl was added to pH 4. Then the mixture was extracted with ether (total volume of 350 25 ml).

The water layer was basified with concentrated ammonia to pH 8-9 and extracted with toluene (total volume of 200 ml). The toluene extract was dried over magnesium sulfate, filtered and evaporated to dryness in vacuo.

30 This crude 1-(6-methoxy-4-quinolyl)-3-[3(R)-ethyl-4(S)-piperidyl]-propanone-2 was dissolved in 80 ml of dimethyl formamide and 6.1 g (61.2 mmol) of propyl iodide in 40 ml of dimethyl formamide was added with stirring.

The reaction mixture was heated to 50-60°C and stirred 35 for 2h.

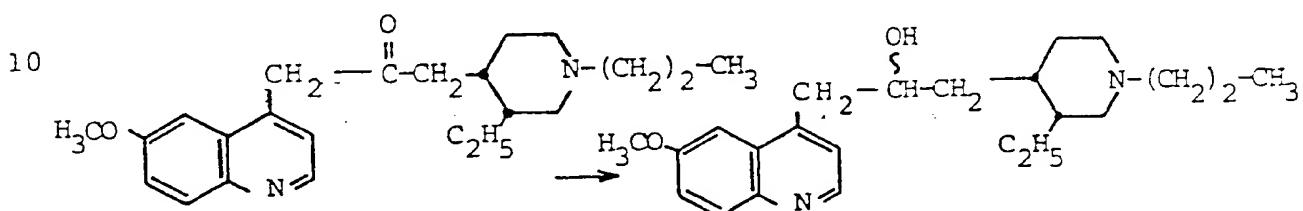
The reaction mixture was poured into 500 ml of water and extracted with toluene (total volume of 450 ml). The toluene extract was dried over magnesium sulfate, filtered

and evaporated to dryness in vacuo. The crude product was purified over Silica Gel G with ethylacetate containing 5% diethylamine as eluent.

The 1-(6-methoxy-4-quinolyl)-3- β (R)-ethyl-N-propyl-4(S)-piperidyl-propanone-2 was obtained as an oil.

Example 97

1-(6-methoxy-4-quinolyl)-3- β (R)-ethyl-N-propyl-4(S)-piperidyl-propanol-2.

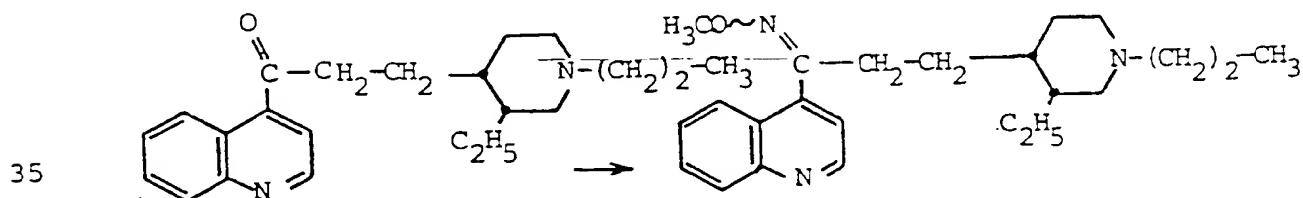


A mixture of 9.0 g (24.5 mmol) 1-(6-methoxy-4-quinolyl)-3- β (R)-ethyl-N-propyl-4(S)-piperidyl-propanone-2 and 0.6 g (15.9 mmol) of sodiumborohydride in 75 ml of methanol containing 0.5 ml of a 4 n sodiumhydroxide solution and 9 ml of water was stirred at room temperature for 3 h. Then 150 ml of water was added and the mixture was extracted with chloroform (total volume of 300 ml). The chloroform extract was dried over magnesium sulfate, filtered and evaporated to dryness in vacuo.

The crude product was purified over Silica Gel G with ethylacetate containing 5% diethylamine as eluent.

The 1-(6-methoxy-4-quinolyl)-3- β (R)-ethyl-N-propyl-4(S)-piperidyl-propanol-2 (mixture of isomers) was obtained as an oil. Melting point of hydrobromide salt is 213°C.

Example 98
30 N-propyl-hydrocinchonicine-O-methyl oxime bioxalate



A mixture of 23.0 g (0.068 mol) of N-propyl-hydrocinchonicine (25) and 8.0 g (0.096 mol) of methoxylamine in 200

ml of absolute ethanol was refluxed for 16 h.

After cooling to room temperature the reaction mixture was evaporated to dryness in vacuo. Water was added to the residue, the mixture basified with concentrated ammonia and 5 extracted with chloroform. The chloroform extract was dried over magnesium sulfate, filtered and evaporated to dryness in vacuo.

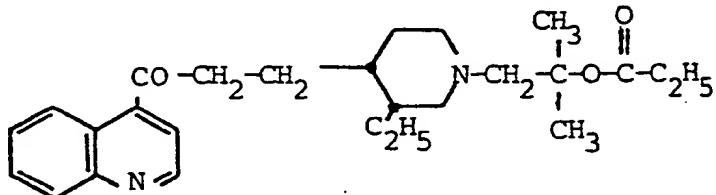
The crude product was purified over Silica Gel G with ethylacetate / diethylamine (50:1) as eluent. The N-propyl-10 hydrocinchonicine-O-methyl oxime (mixture of isomers) was obtained as an oil.

With oxalic acid the base was converted to the oxalate (mol. ratio 1:1.25) with a melting point of 154°C.

Example 99

15 N-(2-methyl-2-propionyloxy)propyl-hydrocinchonicine bifumarate

20



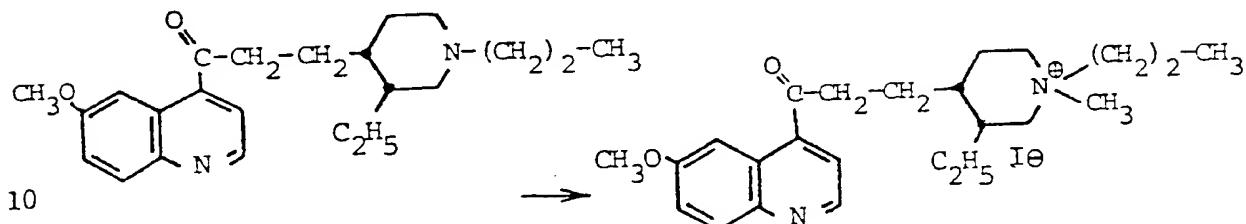
A mixture of 15.0 g (40 mmole) of N-(2-hydroxy-2-methyl) 25 propylhydrocinchonicine, 4.4 g (48 mmole) of propionyl chloride and 5.7 g (56 mmole) of triethyl amine in 300 ml of carbon tetrachloride was refluxed overnight with stirring. The reaction mixture was cooled to room temperature and evaporated to dryness in vacuo. The residue was dissolved in 100 ml of toluene and subsequently extracted with 25 ml of 30 water, 25 ml of a saturated solution of sodium bicarbonate and 25 ml of water. The layers were separated and the organic layer was dried over magnesium sulphate, filtered and concentrated in vacuo.

The residue was dissolved in 50 ml of cyclohexane and 35 chromatographed over a silica gel column with cyclohexane / acetone 3:1 as the eluent. The fraction containing the desired product was evaporated to dryness in vacuo, then the base was dissolved in 25 ml of acetone. To the solution 1.6 g of fuma-

ric acid in isopropylalcohol was added. The salt obtained was isolated and recrystallized from methanol/ethyl acetate. Melting point 132-134°C.

Example 100

5 N-(n-propyl)-hydroquinicine-methiodide.



10 grams (27 mmol) of N-(n-propyl)-hydroquinicine was dissolved in 150 ml of 96% ethylalcohol, to which mixture 5 grams (35 mmol) of methyliodide were added.

15 The mixture was gently heated under reflux until all of the starting material had been consumed; the reaction was followed by means of TLC.

20 After completion of the reaction the reaction mixture was evaporated to dryness in vacuo and recrystallized from an ethylalcohol-water mixture, containing 30% of water. The desired methiodide was obtained after filtration and drying in vacuo. The melting point of N-(n-propyl)-hydroquinicine-methiodide after recrystallisation was 180-185°C.

25 PHARMACOLOGY

Experiment 1 - Effectiveness of the compounds of Examples in spontaneously hypertensive rats

30 Systolic blood pressures were recorded by a modification of the tail cuff method described by I.M. Claxton et al, Eur. J.Pharmacology 37, 179 (1976). An oscilloscope or W+W BP recorder, model 8002, was used to display pulses.

35 Prior to all measurements rats were placed in a heated environment ($33.5 \pm 0.5^\circ\text{C}$) before transfer to a restraining cage. Each determination of blood pressure was the mean of at least 6 readings.

Spontaneously hypertensive rats (aged 12-18 weeks) with systolic blood pressures >170 mm Hg were considered hypertensive. Groups of 6 animals ($n=6$) were used unless specified.

In the following Table the results with certain compounds of the invention, which have been carried out with the above-described method, are mentioned. The numbers of the compounds correspond with those of the Examples.

5

Table 1

Compound No.	dosage mg/kg	change of systolic blood pressures (%) in different time intervals (h)					
		1	2	4	6	24	
10	2	100	-20	-27	-17	-15	+3
	4	10	-34	-25	-23	-21	-4
	6	10	0	-2	-14	-15	-7
	7	10	-26	-27	-36	-28	+4
	8	100	-27	-40	-39	-10	-4
15	11	10	0	1	-1	-6	-
	12	100	-28	-38	-23	0	0
	14	100	-10	-9	-3	1	-1
	16	10	-28	-30	-38	-30	-7
	17	10	-38	-35	-34	-32	+1
20	18	1	-6	-11	-14	-9	+17
	19	10	-8	-17	-23	-30	9
	20	10	-37	-34	-33	-27	0
	21	10	-41	-24	-26	-25	-13
	22	1	-13	-11	-12	-11	-
25	24	10	-5	-16	-14	-15	-5
	25	100	-12	-24	-28	-25	+1
	28	10	3	-5*	-4	-19*	-1
	29	10	-8	-10	-12	-16	+2
	30	100	-23	-34	-35	-48	-17
30	31	10	-3	-2	-11	-15	-3
	34	100	-24	-37	-29	-48	-2
	41	100	-17	-34	-37	-34	-3
	42	100	-38	-35	-40	-25	+3
	44	100	-25	-19	-18	-29	-2
35	60	10	-4	-11	-8	-13	-
	66	100	-7	-13	-20	-13	-3
	67	10	-11	-13	-14	-20	-5 (n=5)

Tabel 1 (cont.)

Compound No.	dosage mg/kg	change of systolic blood pressures (%) in different time intervals (h)					(n=4)
		1	2	4	6	24	
5	68	10	-9	3	-5	-2	2 (n=4)
	70	10	+1	+2	-3	-11	+2
	71	10	-3	-6	-8	-5	+8
	80	100	-17	-28	-44	-44	-7
	82	100	-22	-14	-23	-12	+4
10	83	10	0	-9	-12	-10	12
	84	10	-6	-28	-10	-19	+2
	86	100	+2	-7	-2	-13	+8
	88	100	-9	-15	-23	-9	+14
	89	10	-1	-7	-14	-14	-5
15	90	10	-3	-5	-7	-9	-11
	92	10	-1	-5	-22	-23*	-3 (n=5)
	93	100	-11	-14	-17	-20	3
	94	10	-9	-11	-12	-16	0
	95	100	-16	-23	-35	-23	+5

20

* No pulse in 2 rats

Experiment 2 - Effectiveness of the compounds of Examples in the Guinea Pigs Electrostimulation Test

25 Arrhythmias were induced in guinea pigs by electrostimulation of the right ventricle of the heart. The animals were anaesthetized with urethane (1.2 g/kg i.p.) and artificially respirated before a needle electrode was inserted in the right ventricle of the heart. Substances were given intraduodenally 30 min. before the stimulation at a dose of 32 mg/kg.

The voltage needed for induction of extra systoles in control animals (n=6) was compared with that required for induction of arrhythmias in treated animals (n=6).

35 This method is based on the work of L. Szekeres and G.J. Papp, Naunyn-Schmiedebergs Arch. Exp. Path. Pharmak., 245, 70 (1963).

In the Table the results of certain compounds of the

0030044

invention are mentioned, which have been carried out according to the method described above.

The numbers of the compounds correspond with those of the Examples.

5

Tabel 2

Guinea Pig Electrostimulation Test at 32 mg/kg unless
Indicated in Brackets after Result

<u>Compound Number</u>		<u>Percent Increase in Voltage (%) Required for Arrhythmia</u>
10	2	46
	4	5
	6	Ia*
	7	9
	11	12
15	24	Ia*
	25	48
	26	11
	27	6 (16)
	30	25
20	31	141
	32	121
	35	20 (16)
	38	20
	40	85
25	41	30
	42	28
	45	72
	53	15 (16)
	57	9
30	66	18
	67	Ia*
	68	15
	70	Ia*
	71	23
35	86	56
	89	9
	92	Ia*
	98	58

Table 2 (cont.)

<u>Compound Number</u>	<u>Percent Increase in Voltage (%) Required for Arrhythmia</u>
100	51 (16)

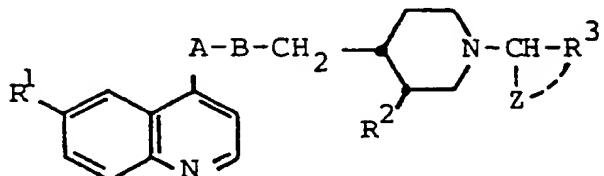
5 * Inactive at the tested dosage.

Some compounds also showed anti depressant activity and some compounds also showed anti-inflammatory activity, in preliminary tests.

C L A I M S

1. A compound of formula 1 or a salt thereof,

5



10 in which A-B is $-\text{CH}_2-\text{CH}_2-$, $-\text{CHOH}-\text{CH}_2-$, $-\text{CH}_2-\text{CHOH}-$, $-\text{C}(\text{O})-\text{CH}_2-$,
 $-\text{CH}_2-\text{C}(\text{O})-$, $-\text{C}(\text{NOR}^4)-\text{CH}_2-$ or $-\text{CH}_2-\text{C}(\text{NOR}^4)-$,

R¹ is hydrogen, hydroxy or lower alkoxy,

R² is ethyl or vinyl,

R³ is C₂₋₈ alkyl, C₁₋₈ hydroxyalkyl, lower alkoxy-

15

alkyl or lower alkanoyloxyalkyl, C₃₋₆ cyclo-

alkyl, hydroxycycloalkyl, lower alkoxyxcyclo-

alkyl or lower alkanoyloxycycloalkyl, cyclo-

alkyl lower alkyl, hydroxy-, lower alkoxy-

or lower alkanoyloxyxcycloalkyl lower alkyl;

20

cyano, cyano lower alkyl, lower alkenyl,

lower alkynyl, tetrahydrofuryl, mono- or

di-lower alkylamino lower alkyl, mono- or

di-lower alkylamino lower hydroxy alkyl;

optionally substituted phenyl, phenyl lower

alkyl or phenyl hydroxy lower alkyl, optional-

ly substituted diphenyl lower alkyl, optional-

ly substituted phenyl lower alkenyl, optionally

substituted benzoyl or benzoyl lower alkyl, optionally

substituted heteroaryl or heteroaryl lower

25

alkyl, or optionally substituted heteroaroyl

or heteroaroyl lower alkyl,

30

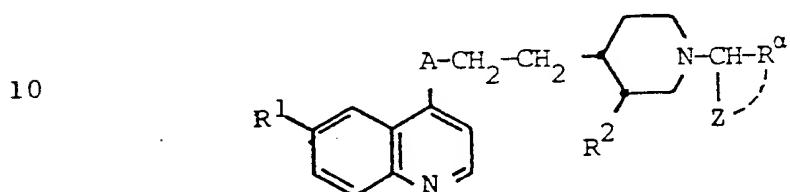
R⁴ is lower alkyl, and

35

Z is hydrogen, lower alkyl or optionally substituted phenyl, or Z and R³ together with the carbon atom to which they are attached form a C₃₋₆ cycloalkyl group,

whereby the substituents at the 3- and 4-position of the piperidine ring are in the cis-position, excluding N- \bar{C}_{2-6} alkyl, C_{2-6} hydroxyalkyl, NN-di-lower alkylamino lower alkyl, optionally substituted C_{7-11} aralkyl substituted derivatives
5 of quinicine and cinchonicine.

2. A compound according to claim 1 or a salt thereof,
as represented by formula 1a,



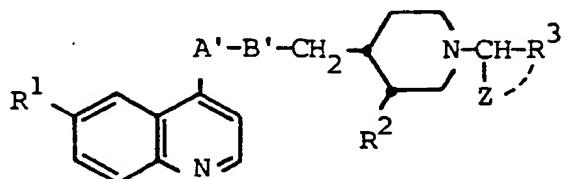
in which A is $-\text{CH}_2-$, $-\text{CHOH}-$ or $-\text{CO}-$,
15 R^1 is hydrogen, hydroxy or lower alkoxy,
 R^2 is ethyl or vinyl,
 R^3 is C_{2-8} alkyl, C_{1-8} hydroxyalkyl, lower alkoxy-
 alkyl or lower alkanoyloxyalkyl, C_{3-6} cycloalkyl, hy-
 droxycycloalkyl, lower alkoxycycloalkyl or lower alka-
 noyloxycycloalkyl, cycloalkyl lower alkyl, hydroxy-,
20 lower alkoxy- or lower alkanoyloxycycloalkyl lower
 alkyl; cyano, cyano lower alkyl, lower alkenyl, lower
 alkynyl, tetrahydrofuryl, mono- or di-lower alkyl-
 amino lower alkyl; mono- or di-lower alkylamino hy-
 droxy lower alkyl; optionally substituted phenyl,
 phenyl lower alkyl or phenyl hydroxy lower alkyl;
 optionally substituted diphenyl lower alkyl, op-
 tionally substituted phenyl lower alkenyl, optional-
 ly substituted benzoyl or benzoyl lower alkyl,
 25 optionally substituted heteroaryl or heteroaryl lower
 alkyl, or optionally substituted heteroaroyl or he-
 roaroyl lower alkyl, and

 Z is hydrogen, lower alkyl or optionally substi-
 tuted phenyl, or Z and R^3 together with the
30 carbon atom to which they are attached form
 a C_{3-6} cycloalkyl group,

excluding the quinicine and cinchonicine derivates as in claim 1.

3. A compound according to claim 1 or a salt thereof, as represented by formula 1b,

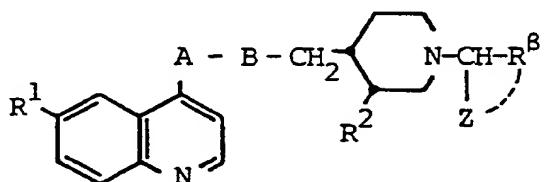
5



in which R¹, R², R³, R⁴ and Z are as defined in claim 1 and
10 A'-B' is -CH₂-CH₂-, -CHOH-CH₂-, -CH₂-CHOH-, -CH₂-C(O)-,
C(NOR⁴)-CH₂- or -CH₂-C(NOR⁴)-.

4. A compound according to claim 1 or a salt thereof, as represented by formula 1c,

15

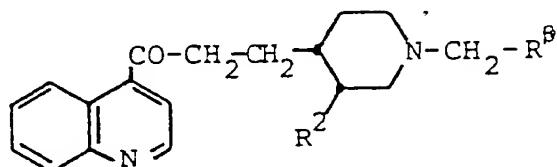


20 in which A-B, R¹, R² and Z are as defined in claim 1 and
R^β is C₁₋₈ lower alkoxyalkyl or lower alkanoyloxy-
alkyl, C₃₋₆ cycloalkyl, hydroxycycloalkyl,
lower alkoxycycloalkyl or lower alkanoyloxy-
cycloalkyl, cycloalkyl lower alkyl, hydroxy-,
25 lower alkoxy- or lower alkanoyloxcycloalkyl
lower alkyl; cyano, cyano lower alkyl, lower
alkenyl, lower alkynyl, tetrahydrofuryl,
mono- or di-lower alkylamino lower hydroxy-
alkyl; optionally substituted phenylhydroxy
30 lower alkyl, optionally substituted diphenyl
lower alkyl, optionally substituted phenyl
lower alkenyl, optionally substituted benzoyl
or benzoyl lower alkyl, optionally substituted
heteroaryl or heteroaryl lower alkyl, or
optionally substituted heteroaroyl or hetero-
35 aroyl lower alkyl, or

β and R^B together with the carbon atom to which they are attached form a C_{3-6} cycloalkyl group.

5. A compound according to claim 1 or a salt thereof, as represented by formula 1d,

5

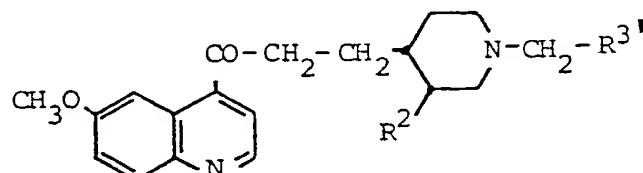


10

in which R^2 and R^B are as defined in claim 4.

6. A compound according to claim 1 or a salt thereof, as represented by formula 1e,

15



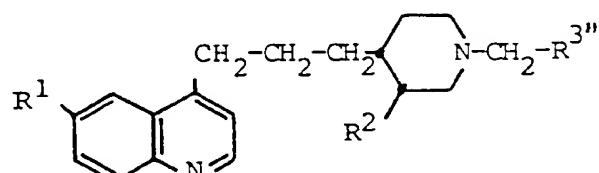
20

in which R^2 is as defined in claim 1, $R^{3'}$ is ethyl, n-butyl, iso-butyl, n-pentyl, iso-pentyl, lower alkoxy lower alkyl, C_{3-6} cycloalkyl lower alkyl, lower alkenyl, optionally substituted phenyl lower alkyl or optionally substituted benzoyl or benzoyl lower alkyl. R^2 is preferably vinyl. Especially preferred are the compounds of formula 1e in which R^2 is vinyl and $R^{3'}$ is 2-benzoylethyl, 2-(4-fluorobenzoyl)ethyl or 1-hydroxy-1-methyl-ethyl.

25

7. A compound according to claim 1 or a salt thereof, as represented by formula 1f,

30



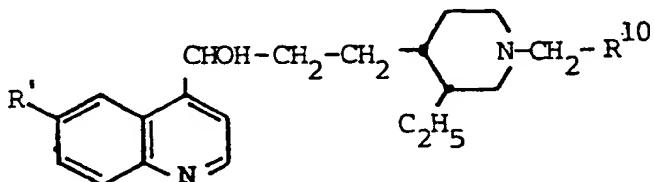
35

in which R^1 and R^2 are as defined in claim 1 and $R^{3''}$ is ethyl, n-butyl, iso-butyl, n-pentyl, iso-pentyl, cyclobutyl,

lower alkenyl, optionally substituted phenyl lower alkyl or optionally substituted benzoyl or benzoyl lower alkyl.

8. A compound according to claim 1 or a salt thereof, as represented by formula 1g,

5

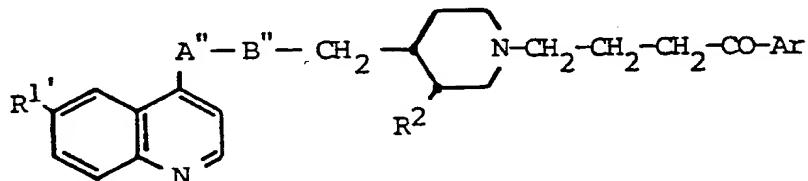


10

in which R¹ is as defined in claim 1 and R¹⁰ is alkyl or -(CH₂)_n-C(O)R¹¹, wherein R¹¹ is an optionally substituted phenyl group and n is 0, 1, 2, 3 or 4.

9. A compound according to claim 1 or a pharmaceutically acceptable salt thereof, as represented by formula 1h,

20

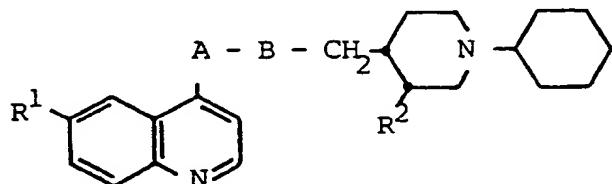


25

in which A''-B'' is -CHOH-CH₂- or -CH₂-CH₂-, R¹ is hydrogen or methoxy, R² is ethyl or vinyl and Ar is phenyl, thiienyl or phenyl substituted by one, two or three groups, selected from fluorine, chlorine or methoxy.

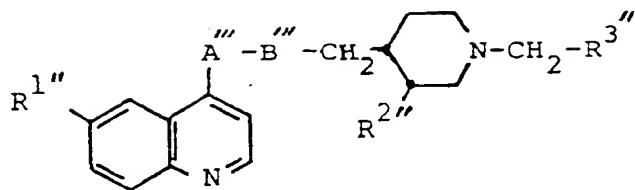
10. A compound according to claim 1 or a pharmaceutically acceptable salt thereof, as represented by formula 1i,

30



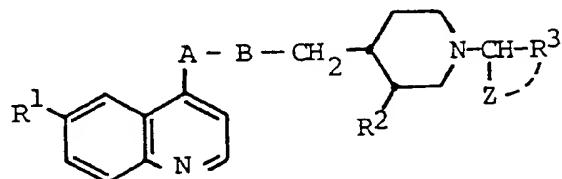
in which A-B, R¹ and R² are as defined in claim 1.

35 11. A compound according to claim 1 or a pharmaceutically acceptable salt thereof, as represented by formula 1j,



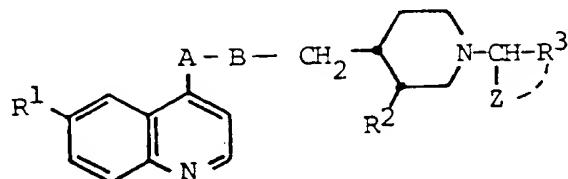
in which A'''-B''' is -CHOH-CH₂- or -CH₂-CHOH- or -CH₂-CH₂-,
R^{1''} is hydrogen or methoxy, R^{2''} is ethyl or vinyl and
R^{3''} is alkyl, hydroxyalkyl or alkoxyalkyl.

12. N-(3-benzoyl)propyl-hydrocinchonicine.
13. N-(3-benzoyl)propyl-hydrocinchonicinol-1.
14. N-(3-benzoyl)propyl-desoxo-hydrocinchonicine.
15. N-propyl-hydroquinicinol-1.
16. N-butyl-hydroquinicinol-1.
17. N-pentyl-hydroquinicinol-1.
18. N-pentyl-hydrocinchonicinol-1.
19. N-(4-methoxy)butyl-hydrocinchonicine.
20. A method for the preparation of a compound of formula 1,



as defined in claim 1, characterized in that such compound is prepared in a manner known per se for the synthesis of analogous compounds.

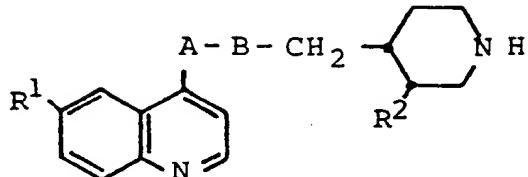
21. A process for the preparation of a compound of formula 1,



as claimed in claim 1 or a salt thereof, characterized in that

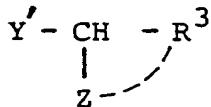
A) a compound of formula 5

5



10 in which A-B, R¹ and R² are as defined in claim 1, is alkylated with a compound of formula 6,

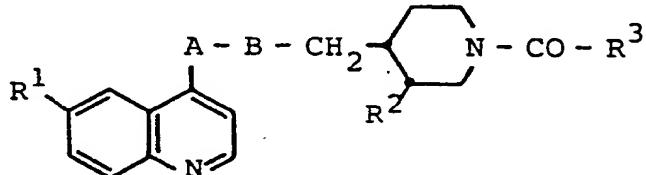
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in which R³ and Z are as defined in claim 1 and Y' is a nucleophilic leaving group, particularly chlorine, bromine, 20 iodine, aryl-, aralkyl- or alkylsulphonyloxy, and especially mesyloxy or tosyloxy,

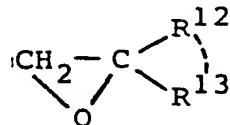
or B) a compound of formula 7,

25



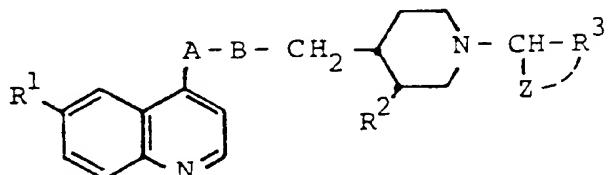
in which A-B, R¹ and R² are as defined in claim 1, is reduced, 30 to give a compound of formula 1 in which Z is hydrogen, or C) a compound of formula 5 above is reacted with an epoxide of formula 8,

35



in which R¹² is C₁₋₆ alkyl, lower alkenyl, lower alkynyl, optionally substituted phenyl and phenyl lower alkyl, and R¹³ is hydrogen and lower alkyl or R¹² and R¹³, together with the carbon atom to which they are attached, form a C₃₋₆ cycloalkyl group, to give a compound of formula 1,

10



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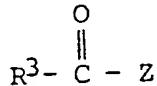
in which Z is hydrogen and R³ is as defined above,
or D) a compound of formula 1 in which R³ contains a hydroxy group is alkylated,

15

or E) a compound of formula 1 in which R³ contains a hydroxy group is acylated,

or F) a compound of formula 5 above is reacted with a compound of formula 9,

20



25

in which R³ and Z are as defined above, in the presence of a reducing agent,

30

or G) a compound of formula 1, in which A-B contains a carbonyl group is reacted with an O-substituted hydroxyl-amine derivative of formula YO-NH₂, in which Y is R⁴ or a group replaceable by or convertable into R⁴, R⁴ being as previously defined, whereafter the resulting compound in which Y ≠ R⁴ is converted in a compound in which Y = R⁴,

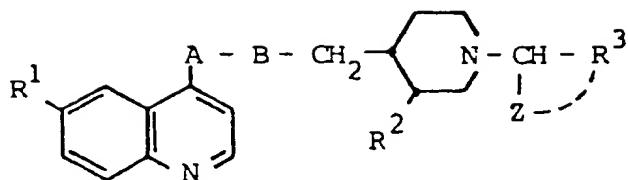
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or H) a compound of formula 1 in which A-B is -CO-CH₂- or -CH₂-CO- is partially or completely reduced to -CHOH-CH₂- or -CH₂-CHOH-, or -CH₂-CH₂, respectively.

22. A method for the preparation of a pharmaceutical composition, characterized in that at least a compound of

formula 1,

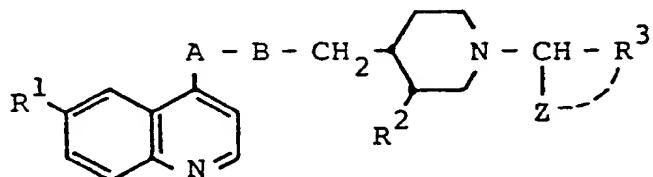
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as defined in claim 1, or a pharmaceutically acceptable salt thereof, is brought in a form suitable for therapeutic purposes.

23. A method of treating mammals, in particular humans, suffering from e.g. cardiovascular diseases, which comprises administering an effective amount of a compound of formula 1,

15



20 as defined hereinbefore or a pharmaceutically acceptable salt thereof, preferably in the form of a pharmaceutical composition.

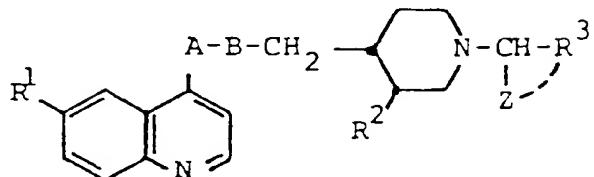
24. Compounds, pharmaceutical compositions and methods, as described in the specification and examples.

25. 25. A compound of the formula 1 or a salt thereof, for use in treating cardiovascular disorders.

CLAIMS FOR AUSTRIA

1. A process for the preparation of novel quinoline derivatives characterized in that one prepares compounds of formula 1 or salts thereof,

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in which A-B is $-\text{CH}_2-\text{CH}_2-$, $-\text{CHOH}-\text{CH}_2-$, $-\text{CH}_2-\text{CHOH}-$, $-\text{C}(\text{O})-\text{CH}_2-$,
 $-\text{CH}_2-\text{C}(\text{O})-$, $-\text{C}(\text{NOR}^4)-\text{CH}_2-$ or $-\text{CH}_2-\text{C}(\text{NOR}^4)-$,

R^1 is hydrogen, hydroxy or lower alkoxy,

R^2 is ethyl or vinyl,

15 R^3 is C_{2-8} alkyl, C_{1-8} hydroxyalkyl, lower alkoxy-alkyl or lower alkanoyloxyalkyl, C_{3-6} cycloalkyl, hydroxycycloalkyl, lower alkoxyxycycloalkyl or lower alkanoyloxycycloalkyl, cycloalkyl lower alkyl, hydroxy-, lower alkoxy- or lower alkanoyloxycycloalkyl lower alkyl; cyano, cyano lower alkyl, lower alkenyl, lower alkynyl, tetrahydrofuryl, mono- or di-lower alkylamino lower alkyl, mono- or di-lower alkylamino lower hydroxy alkyl;

20

25 optionally substituted phenyl, phenyl lower alkyl or phenyl hydroxy lower alkyl, optionally substituted diphenyl lower alkyl, optionally substituted phenyl lower alkenyl, optionally substituted benzoyl or benzoyl lower alkyl, optionally substituted heteroaryl or heteroaryl lower alkyl, or optionally substituted heteroaroyl or heteroaroyl lower alkyl,

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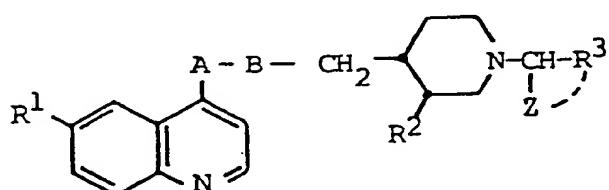
R^4 is lower alkyl, and

35 Z is hydrogen, lower alkyl or optionally substituted phenyl, or Z and R^3 together with the carbon atom to which they are attached form a C_{3-6} cycloalkyl group,

whereby the substituents at the 3- and 5-position of the piperidine ring are in the cis-position, excluding N- \overline{C}_{2-6} alkyl, C_{2-6} hydroxyalkyl, NN-di-lower alkylamino lower alkyl, optionally substituted C_{7-11} aralkyl substituted derivatives 5 of quinicine and cinchonicine, in a manner known per se for the synthesis of analogous compounds.

2. A process for the preparation of a compound of formula 1,

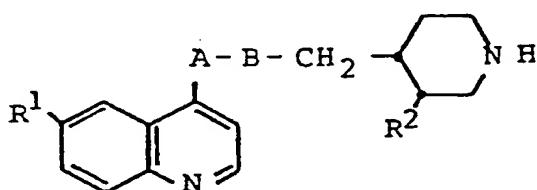
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as claimed in claim 1 or a salt thereof, characterized in that
A) a compound of formula 5

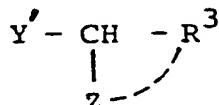
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in which A-B, R¹ and R² are as defined in claim 1, is alkylated with a compound of formula 6,

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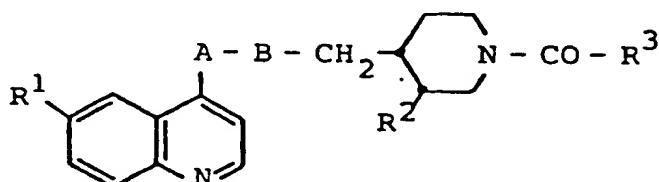


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in which R³ and Z are as defined in claim 1 and Y' is a nucleophilic leaving group, particularly chlorine, bromine, iodine, aryl-, aralkyl- or alkylsulphonyloxy, and especially mesyloxy or tosyloxy,

or B) a compound of formula 7,

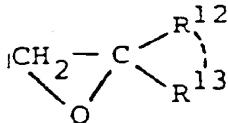
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in which A-B, R¹ and R² are as defined in claim 1, is reduced, to give a compound of formula 1 in which Z is hydrogen,

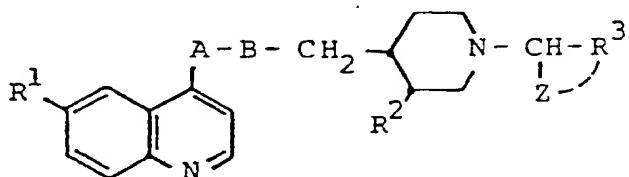
or C) a compound of formula 5 above is reacted with an epoxide of formula 8,

5



10 in which R¹² is C₁₋₆ alkyl, lower alkenyl, lower alkynyl, optionally substituted phenyl and phenyl lower alkyl, and R¹³ is hydrogen and lower alkyl or R¹² and R¹³, together with the carbon atom to which they are attached, form a C₃₋₆ cycloalkyl group, to give a compound of formula 1,

15



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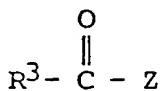
in which Z is hydrogen and R³ is as defined above,

or D) a compound of formula 1 in which R³ contains a hydroxy group is alkylated,

25 or E) a compound of formula 1 in which R³ contains a hydroxy group is acylated,

or F) a compound of formula 5 above is reacted with a compound of formula 9,

30

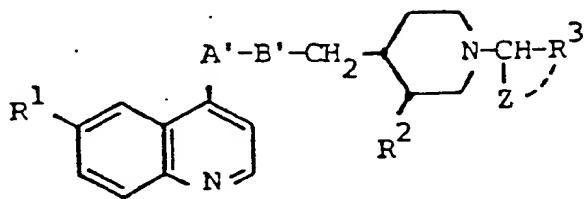


in which R³ and Z are as defined above, in the presence of 35 a reducing agent,

or G) a compound of formula 1, in which A-B contains a carbonyl group is reacted with an O-substituted hydroxyl-amine derivative of formula YO-NH₂, in which Y is R⁴ or a group replaceable by or convertible into R⁴, R⁴ being as 40 previously defined, whereafter the resulting compound is

4. The process according to claim 1 or 2 characterized in that one prepares a compound of formula 1b, or a salt thereof

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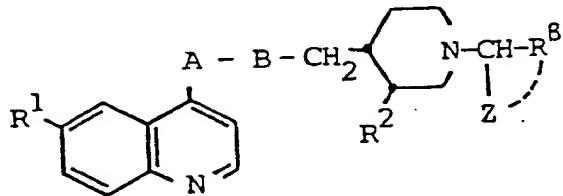


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in which R¹, R², R³, R⁴ and Z are as defined in claim 1 and A'-B' is -CH₂-CH₂-, -CHOH-CH₂-, -CH₂-CHOH-, -CH₂-C(O)-, C(NOR⁴)-CH₂- or -CH₂-C(NOR⁴)-.

5. The process according to claim 1 or 2 characterized
15 in that one prepares a compound of formula 1c, or a salt
thereof

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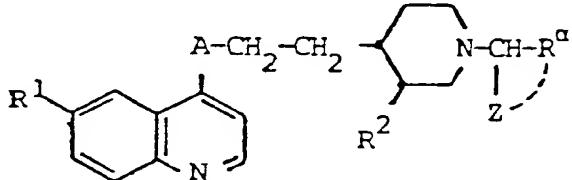
in which A-B, R¹, R² and Z are as defined in claim 1 and
25 R⁸ is C₁₋₈ lower alkoxyalkyl or lower alkanoyloxy-alkyl, C₃₋₆ cycloalkyl, hydroxycycloalkyl, lower alkoxy-cycloalkyl or lower alkanoyloxy-cycloalkyl, cycloalkyl lower alkyl, hydroxy-, lower alkoxy- or lower alkanoyloxy-cycloalkyl lower alkyl; cyano, cyano lower alkyl, lower alkenyl, lower alkynyl, tetrahydrofuryl, mono- or di-lower alkylamino lower hydroxy-alkyl; optionally substituted phenylhydroxy lower alkyl, optionally substituted diphenyl lower alkyl, optionally substituted phenyl lower alkenyl, optionally substituted benzoyl or benzoyl lower alkyl, optionally substituted heteroaryl or heteroaryl lower alkyl, or
35 optionally substituted heteroaroyl or heteroaroyl lower alkyl, or
40

which $Y \neq R^4$ is converted in a compound in which $Y = R^4$,

or H) a compound of formula I in which A-B is $-CO-CH_2-$ or $-CH_2-CO-$ is partially or completely reduced to $-CHOH-CH_2-$ or $-CH_2-CHOH-$, or $-CH_2-CH_2-$, respectively.

5 3. The process according to claim 1 or 2, characterized in that one prepares a compound of formula 1a or a salt thereof,

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15 in which A is $-CH_2-$, $-CHOH-$ or $-CO-$,

R^1 is hydrogen, hydroxy or lower alkoxy,

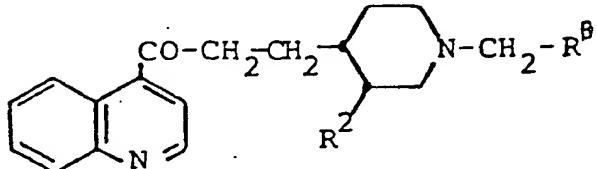
R^2 is ethyl or vinyl,

R^3 is C_{2-8} alkyl, C_{1-8} hydroxyalkyl, lower alkoxyalkyl or lower alkanoyloxyalkyl, C_{3-6} cycloalkyl, hydroxycycloalkyl, lower alkoxycycloalkyl or lower alkanoyloxycycloalkyl, cycloalkyl lower alkyl, hydroxy-, lower alkoxy- or lower alkanoyloxycycloalkyl lower alkyl; cyano, cyano lower alkyl, lower alkenyl, lower alkynyl, tetrahydrofuryl, mono- or di-lower alkyl-amino lower alkyl; mono- or di-lower alkyl-amino hydroxy lower alkyl; optionally substituted phenyl, phenyl lower alkyl or phenyl hydroxy lower alkyl; optionally substituted diphenyl lower alkyl, optionally substituted phenyl lower alkenyl, optionally substituted benzoyl or benzoyl lower alkyl, optionally substituted heteroaryl or heteroaryl lower alkyl, or optionally substituted heteroaroyl or heteroaroyl lower alkyl, and

Z is hydrogen, lower alkyl or optionally substituted phenyl, or Z and R^3 together with the carbon atom to which they are attached form a C_{3-6} cycloalkyl group, excluding the quinicine and cinchonicine derivates as in claim 1.

Z and R^B together with the carbon atom to which they are attached form a C_{3-6} cycloalkyl group.

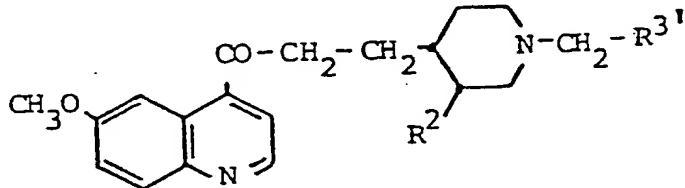
6. The process according to claim 1 or 2 characterized in that one prepares a compound of formula 1d, or
5 a salt thereof,



10

in which R^2 and R^B are as defined in claim 4.

7. The process according to claim 1 or 2 characterized in that one prepares a compound of formula 1e, or a salt thereof,

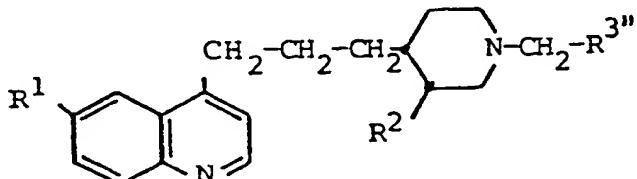


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in which R^2 is as defined in claim 1, R^3' is ethyl, n-butyl, iso-butyl, n-pentyl, iso-pentyl, lower alkoxy lower alkyl, C_{3-6} cycloalkyl lower alkyl, lower alkenyl, optionally substituted phenyl lower alkyl or optionally substituted benzoyl or benzoyl lower alkyl. R^2 is preferably vinyl. Especially preferred are the compounds of formula 1e in which R^2 is vinyl and R^3' is 2-benzoylethyl, 2-(4-fluorobenzoyl)ethyl or 1-hydroxy-1-methyl-ethyl.

8. The process according to claim 1 or 2 characterized in that one prepares a compound of formula 1f, or a salt thereof.

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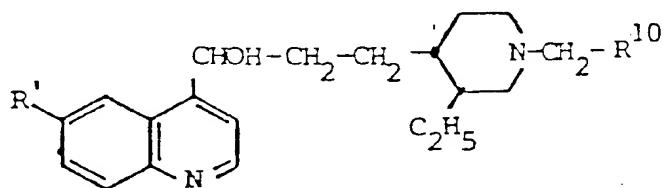


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in which R¹ and R² are as defined in claim 1 and R³ is ethyl, n-butyl, iso-butyl, n-pentyl, iso-pentyl, cyclobutyl, lower alkenyl, optionally substituted phenyl lower alkyl or optionally substituted benzoyl or benzoyl lower alkyl.

5 9. The process according to claim 1 or 2 characterized in that one prepares a compound of formula 1g, or a salt thereof,

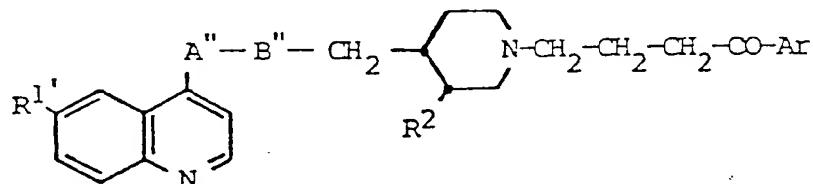
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15 in which R¹ is as defined in claim 1 and R¹⁰ is alkyl or -(CH₂)_n-C(O)R¹¹, wherein R¹¹ is an optionally substituted phenyl group and n is 0, 1, 2, 3 or 4.

10. The process according to claim 1 or 2 characterized in that one prepares a compound of formula 1h, or a
20 salt thereof,

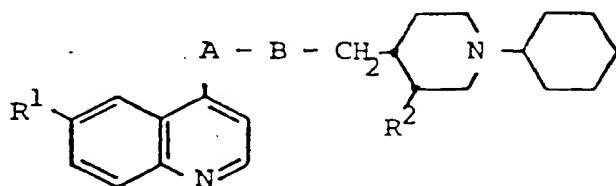
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in which A''-B'' is -CHOH-CH₂- or -CH₂-CH₂-, R¹ is hydrogen or methoxy, R² is ethyl or vinyl and Ar is phenyl, thiophenyl or phenyl substituted by one, two or three groups, selected from fluorine, chlorine or methoxy.

30 11. The process according to claim 1 or 2 characterized in that one prepares a compound of formula 1i, or a
salt thereof

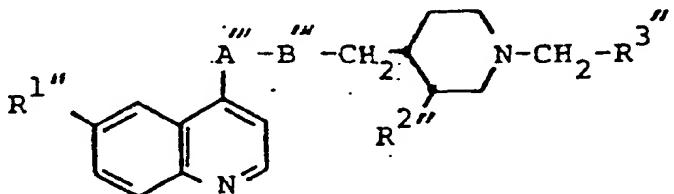
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in which A, R¹ and R² are as defined in claim 1.

12. The process according to claim 1 or 2 characterized in that one prepares a compound of formula 1j, or a salt thereof,

5



10

in which A'''-B''' is -CHOH-CH₂- , -CH₂-CHOH- or -CH₂-CH₂- , R^{1'''} is hydrogen or methoxy, R^{2'''} is ethyl or vinyl and R^{3'''} is alkyl, hydroxyalkyl or alkoxyalkyl.

15 13. The process according to one of the preceding claims 1-12 characterized in that one prepares N-(3-benzoyl)-propyl-hydrocinchonicine.

14. The process according to one of the preceding claims 1-12 characterized in that one prepares N-(3-benzoyl)-propyl-hydrocinchonicinol-1.

15. The process according to one of the preceding claims 1-12 characterized in that one prepares N-(3-benzoyl)-propyl-desoxo-hydrocinchonicine.

16. The process according to one of the preceding 25 claims 1-12 characterized in that one prepares N-propyl-hydroquinicinol-1.

17. The process according to one of the preceding claims 1-12 characterized in that one prepares N-butyl-hydroquinicinol-1.

30 18. The process according to one of the preceding claims 1-12 characterized in that one prepares N-pentyl-hydroquinicinol-1.

19. The process according to one of the preceding 35 claims 1-12 characterized in that one prepares N-pentyl-hydrocinchonicinol-1.

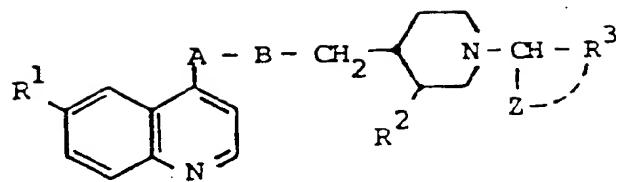
20. The process according to one of the preceding claims 1-12 characterized in that one prepares N-(4-methoxy)butyl-hydrocinchonicine.

40 21. A method for the preparation of a pharmaceutical composition, characterized in that at least a compound of

formula

0030044

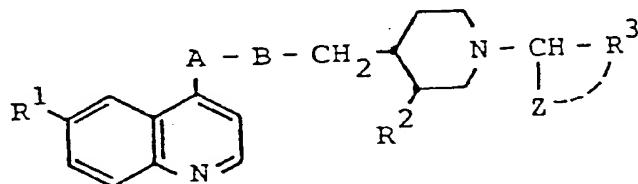
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as defined in claim 1, or a pharmaceutically acceptable
10 salt thereof, is brought in a form suitable for therapeutic
purposes.

22. A method of treating mammals, in particular
humans, suffering from e.g. cardiovascular diseases, which
comprises administering an effective amount of a compound of
15 formula 1,

20



as defined hereinbefore or a pharmaceutically acceptable salt
thereof, preferably in the form of a pharmaceutical composi-
25 tion.

